

77 Rec'd PCT/PTO 26 NOV 2001

FORM PTO-1390 (Rev. 1/98) (adapted)

U.S. DEPARTMENT OF COMMERCE — PATENT AND TRADEMARK OFFICE		ATTORNEY DOCKET NUMBER
TRANSMITTAL LETTER TO THE UNITED STATES		ABR 022 US
DESIGNATED/ELECTED OFFICE (DO/EO/US)		US Appl'n No. (if known, 37 CFR 1.5)
CONCERNING A FILING UNDER 35 U.S.C. 371		09/926596
INTERNATIONAL APPL'N NO.	INTERNATIONAL FILING DATE	PRIORITY DATE CLAIMED
PCT/SE00/01079	26 MAY 2000	28 MAY 1999

Applicant herewith submits to the United States Designated/Elected Office (DO/EO/US) the following items and other information:

1. ☒ This is a FIRST submission of items concerning a filing under 35 U.S.C. 371.
2. ☐ This is a SECOND or SUBSEQUENT submission of items concerning a filing under 35 U.S.C. 371.
3. ☒ This express request to begin national examination procedures (35 U.S.C. 371(f)) at any time rather than delay examination until the expiration of the applicable time limit set in 35 U.S.C. 371(b) and PCT Articles 22 and 39(1).
4. ☒ A proper Demand for International Preliminary Examination was made by the 19th month from the earliest claimed priority date.
5. ☒ A copy of the International Application as filed (35 U.S.C. 371(c)(2)).
 - a. ☒ is transmitted herewith (required only if not transmitted by the International Bureau).
 - b. ☒ has been transmitted by the International Bureau.
 - c. ☐ is not required, as the application was filed in the United States Receiving Office (RO/US).
6. ☐ A translation of the International Application into English (35 U.S.C. 371(c)(2)).
7. ☒ Amendments to the claims of the International Application under PCT Article 19 (35 U.S.C. 371(c)(3)).
 - a. ☒ are transmitted herewith (required only if not transmitted by the International Bureau).
 - b. ☐ have been transmitted by the International Bureau.
 - c. ☐ have not been made and will not be made.
8. ☐ A translation of the amendments to the claims under PCT Article 19 (35 U.S.C. 371(c)(3)).
9. ☒ An oath or declaration of the inventors (unexecuted) (35 U.S.C. 371(c)(4)).
10. ☐ A translation of the annexes to the International Preliminary Examination Report under PCT Article 36 (35 U.S.C. 371(c)(5))

Items 11. to 18. below concern document(s) or information included:

11. ☐ An Information Disclosure Statement under 37 CFR 1.97 and 1.98.
12. ☐ An assignment document for recording. A separate cover sheet in compliance with 37 CFR 3.28 and 3.31 is included.
13. ☒ A FIRST preliminary amendment.
 ☐ A SECOND or SUBSEQUENT preliminary amendment.
14. ☐ A substitute specification.
15. ☐ A change of power of attorney and/or address letter.
16. ☐ Formal drawings
17. ☒ Return receipt transmittal (MPEP §503) (specifically itemized).
18. ☐ Other items of information:

U.S. APPL'N NO. (if known, 37 CFR 1.5) 09/926596	INT'L APPL'N NO. PCT/SE00/01079	ATTORNEY DOCKET NO. ABR 022 US
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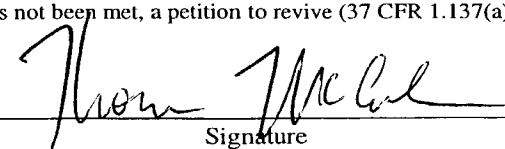
19. <input checked="" type="checkbox"/> The following fees are submitted:				CALCULATIONS		PTO USE ONLY	
BASIC NATIONAL FEE (37 CFR 1.492(a)(1)-(5):							
Neither international preliminary examination fee (37 CFR 1.482) nor international search fee (37 CFR 1.445(a)(2)) paid to USPTO and international Search Report not prepared by the EPO or JPO.....				\$1040.00			
International preliminary examination fee (37 CFR 1.482) not paid to USPTO but International Search Report prepared by the EPO or JPO.....				\$890.00			
International preliminary examination fee (37 CFR 1.482) not paid to USPTO but international search fee paid to USPTO (37 CFR 1.445(a)(2)).....				\$740.00			
International preliminary examination fee (37 CFR 1.482) paid to USPTO but all claims did not satisfy provisions of PCT Article 33(1)-(4).....				\$710.00			
International preliminary examination fee (37 CFR 1.482) paid to USPTO and all claims satisfied provisions of PCT Article 33(1)-(4).....				\$100.00			
ENTER APPROPRIATE BASIC FEE AMOUNT =				\$ 1,040.00			
Surcharge of \$130.00 for furnishing oath or declaration later than <input type="checkbox"/> 20 <input checked="" type="checkbox"/> 30 months from the earliest claimed priority date (37 (CFR 1.492(e)).				\$ 130.00			
CLAIMS	NUMBER FILED	NUMBER EXTRA	RATE				
Total claims	18 - 20 =	0	x \$18.00	\$			
Ind. claims	12 - 3 =	9	x \$84.00	\$ 756.00			
MULTIPLE DEPENDENT CLAIM(s) (if applicable)			+ \$280.00	\$			
TOTAL OF ABOVE CALCULATIONS =				\$			
Reduction of 1/2 for filing by small entity, if applicable. A Small Entity Statement must also be filed (Note 37 CFR 1.9, 1.27, 1.28)				\$			
SUBTOTAL =				\$ 1,926.00			
Processing fee of \$130.00 for furnishing the English translation later than <input type="checkbox"/> 20 <input type="checkbox"/> 30 months from the earliest claimed priority date (37 (CFR 1.492(f)).				\$			
TOTAL NATIONAL FEE =				\$ 1,926.00			
Fee for recording the enclosed assignment (37 CFR 1.21(h)). The assignment must be accompanied by an appropriate cover sheet (37 CFR 3.28, 3.31). \$40.00 per property				+			
TOTAL FEES ENCLOSED =				\$ 1,926.00			
Amount to be:				refunded:		\$	
				charged:		\$	

- a. ☐ A check in the amount of \$ _____ to cover the above fees is enclosed.
- b. ☒ Please charge my Deposit Acct. No. 50-0828 (PowderJect Tech.) in the amount of \$ 1,926.00 to cover the above fees.
- c. ☒ The Commissioner is hereby authorized to charge any additional fees which may be required, or credit any overpayment to Deposit Acct. No. 50-0828 (PowderJect Tech.).

NOTE: Where an appropriate time limit under 37 CFR 1.494 or 1.495 has not been met, a petition to revive (37 CFR 1.137(a) or (b)) must be filed and granted to restore the application to pending status.

SEND ALL CORRESPONDENCE TO:

Thomas P. McCracken
PowderJect Technologies, Inc.
6511 Dumbarton Circle
Fremont, CA 94555


Signature
Thomas P. McCracken, Reg. No. 38,548
Tel. 510-742-9700, ext. 209
Fax 510-742-9720

PTO/PCT Rec'd

08 JUL 2002

Patent

Attorney's Docket No. 033267-011

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

In re Patent Application of

Anders FOLKESSON et al

Application No.: 09/926,596

Filed: November 26, 2001

For: FIMBRIAL PROTEINS



)
)
) Group Art Unit:

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) Examiner:

)
) Confirmation No.: 6671

SUPPLEMENTAL PRELIMINARY AMENDMENT

Assistant Commissioner for Patents
Washington, D.C. 20231

Sir:

Further to the Preliminary Amendment filed on November 26, 2001, and in response to the Notification of Missing Requirements Under 35 U.S.C. 371 mailed on January 7, 2002, please amend the above-identified application as follows:

IN THE SPECIFICATION:

In compliance with 37 C.F.R. § 1.823(a), please substitute the attached paper copy of the Sequence Listing for the originally filed Sequence Listing on pages 14-66 of the application.

Please replace the paragraph at page 4, lines 3-9 with the following paragraph:

--Figure 2: Schematic representation of the pTY52 cosmid comprising the *tcf*-operon

SEQ ID NO:6.

A *tcf* specific PCR fragment of 11105 bp was cloned into the Expand vector I cosmid (Roche). The insert is represented with a thick black line while vector sequences are represented with thin lines. Relevant restriction sites sequences are indicated. The position of the *tcf*-operon, i.e. *tcfA*, *B*, *C* and *D* (SEQ ID NO:6) is represented by a shaded arrow.--

Please replace the paragraph at page 4, lines 15-21 with the following paragraph:

--Sequence listing

SEQ ID NO:1 - DNA sequence of the genes encoding the precursor of the saf fimbriae unit of *Salmonella enterica* subspecies I.

SEQ ID NO:6 - DNA sequence of the genes which encode the precursor of the *tcf* fimbriae unit of *Salmonella enterica* subspecies I serovar Typhi.--

Please replace the paragraph at page 4, lines 23-29 with the following paragraph:

--Deposit information

The phages carrying the inserted SEQ ID NO:1, i.e. phages clones B1, D1, F11 and N10 (see Figure 1) have been given the ECACC Accession numbers 99051922, 99051923, 99051924, and 99051925, respectively.

The cosmid carrying the inserted SEQ ID NO:6, i.e. cosmid pTY52 (see Figure 2) has been given the ECACC Accession number 99051926.--

IN THE CLAIMS:

Please amend the claims as follows (all pending claims are reproduced for the Examiner's convenience):

1. (Amended) A peptide encoded by a nucleotide sequence selected from SEQ ID NO:1 and SEQ ID NO:6 for use in medicine.
2. (Amended) Antibodies directed against a peptide encoded by a nucleotide sequence selected from SEQ ID NO:1 and SEQ ID NO:6 for use in medicine.
3. (Amended) A nucleotide sequence selected from SEQ ID NO:1 and SEQ ID NO:6 for use in medicine.
4. (Amended) A vaccine for the protection against diseases caused by *Salmonella enterica* subspecies I, comprising a peptide encoded by a nucleotide sequence selected from SEQ ID NO:1, or antibodies directed against said peptide and, optionally, a pharmaceutically acceptable carrier.
5. (Amended) A vaccine for the protection against diseases caused by *Salmonella enterica* subspecies I serovar Typhi, comprising a peptide encoded by a nucleotide sequence selected from SEQ ID NO:6 or antibodies directed against said peptide and, optionally, a pharmaceutically acceptable carrier.

6. (Amended) A nucleic acid vaccine for the protection against diseases caused by *Salmonella enterica* subspecies I, comprising a nucleic acid sequence selected from SEQ ID NO:1 and, optionally, a pharmaceutically acceptable carrier.

7. (Amended) A nucleic acid vaccine for the protection against diseases caused by *Salmonella enterica* subspecies I serovar Typhi, comprising a nucleic acid sequence selected from SEQ ID NO:6 and, optionally, a pharmaceutically acceptable carrier.

8. (Amended) A vector vaccine for the protection against diseases caused by *Salmonella enterica* subspecies I, comprising a host in which a recombinant vector comprising a nucleic acid sequence selected from SEQ ID NO:1, has been inserted and, optionally, a pharmaceutically acceptable carrier.

9. (Amended) A vector vaccine for the protection against diseases caused by *Salmonella enterica* subspecies I serovar Typhi, comprising a host in which a recombinant vector comprising a nucleic acid sequence selected from SEQ ID NO:6, has been inserted and, optionally, a pharmaceutically acceptable carrier.

10. A method for vaccinating a mammal against diseases caused by *Salmonella enterica* subspecies I, comprising administering a vaccine according to claim 4 to said mammal.

11. A method for vaccinating a mammal against diseases caused by *Salmonella enterica* subspecies I serovar Typhi, comprising administering a vaccine according to claim 5 to said mammal.

12. (Amended) Antibodies directed against a peptide encoded by a nucleotide sequence selected from SEQ ID NO:1, and from SEQ ID NO:6, for use in a diagnostic method.

13. (Amended) Peptide encoded by a nucleotide sequence selected from SEQ ID NO:1, and SEQ ID NO:6, for use in a diagnostic method.

14. (Amended) Primers for, or probes that hybridize with, a nucleotide sequence selected from SEQ ID NO:1 and SEQ ID NO:6, for use in a diagnostic method for the purpose of detecting *Salmonella enterica* subspecies I.

15. A method for vaccinating a mammal against diseases caused by *Salmonella enterica* subspecies I, comprising administering a vaccine according to claim 6 to said mammal.

16. A method for vaccinating a mammal against diseases caused by *Salmonella enterica* subspecies I, comprising administering a vaccine according to claim 8 to said mammal.

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17. A method for vaccinating a mammal against diseases caused by *Salmonella enterica* subspecies I serovar Typhi, comprising administering a vaccine according to claim 7 to said mammal.

18. A method for vaccinating a mammal against diseases caused by *Salmonella enterica* subspecies I, comprising administering a vaccine according to claim 9 to said mammal.

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REMARKS

Entry of the foregoing and prompt and favorable consideration of the subject application, in light of the following remarks, are respectfully requested.

By the foregoing Supplemental Preliminary Amendment, the specification has been amended to replace the originally filed Sequence Listing with the attached copy of the Sequence Listing, which was filed in parent Application No. PCT/SE00/01079. Although the presently filed Sequence Listing is believed to be contain identical material to combined Sequence Listings 1 and 2 as originally filed with the present specification, this submission is being made in accordance with the statement pursuant to 37 C.F.R. § 1.821(f), which is submitted herewith. No new matter is believed to have been added.

In addition, all references to SEQ ID NO:2 in the specification and claims have been amended to recite "SEQ ID NO:6". The basis for this amendment is the following: The specification as originally filed was filed with two sequence listings, Sequence Listing No. 1, containing SEQ I NOS:1-5, and Sequence Listing No. 2, containing an additional SEQ ID NOS:1-6 (i.e., which were not duplicative of SEQ ID NOS:1-5 of Sequence Listing No. 1).

In the application as filed, sequence number 1 of Sequence Listing number 2 was referred to as SEQ ID NO 2. That only sequence no. 1 of Sequence Listing no. 2 could have been intended by this indication is clear from a comparison of *page 4, line 4* of the description , where SEQ ID NO 2 is identified as the tcf-operon, versus the information given on *page 36 of the Sequence Listing part under items <223>* referring to sequence no. 1 of Sequence Listing no. 2. Namely, under these items, it is stated that the sequence

no. 1 of Sequence Listing no. 2 is the nucleic sequence of the tcf A, B, C and D putative fimbrial subunits respectively and of the tinR putative transcriptional regulator, i.e., is the sequence of the tcf operon. Therefore, it would have been obvious to anyone studying the application as filed that SEQ ID NO:2 referred to sequence no. 1 of Sequence Listing no. 2. Due to renumbering of the sequences, as indicated herein above, this is now sequence number 6, and should consequently be referred to as SEQ ID NO:6.

To further demonstrate that only this sequence could have been intended, it may be noted that in Sequence Listing no. 2 of the application as filed, only the first sequence was a nucleotide acid sequence, the others being protein sequences.

Moreover, on page 4, lines 16-21, the sequences referred to are DNA sequences. The only DNA sequences appearing in the sequence listing part are present SEQ ID NO:1 and SEQ ID NO:6, since all the other sequences are protein sequences. A comparison of the description of the two sequences on page 4, lines 16-21 versus the entries made under items <223> on page 1 and 36, respectively of the Sequence Listing part clearly shows that the sequence described on page 4, lines 16-18 is SEQ ID NO:1 and the sequence described on page 4, lines 19-21 is SEQ ID NO:6.

In the event that there are any questions relating to this Supplemental Preliminary Amendment, or the application in general, it would be appreciated if the Examiner would telephone the undersigned attorney concerning such questions so that prosecution of this application may be expedited.

Respectfully submitted,

BURNS, DOANE, SWECKER & MATHIS, L.L.P.

By: Sharon E. Crane
Sharon E. Crane, Ph.D.
Registration No. 36,113

P.O. Box 1404
Alexandria, Virginia 22313-1404
(703) 836-6620

Date: July 8, 2002

Attachment to Supplemental Preliminary Amendment filed July 8, 2002

Mark-up of Specification

Paragraph at page 4, lines 3-9

Figure 2: Schematic representation of the pTY52 cosmid comprising the *tcf*-operon
 [(SEQ ID NO 2)] SEQ ID NO:6.

A *tcf* specific PCR fragment of 11105 bp was cloned into the Expand vector I cosmid (Roche). The insert is represented with a thick black line while vector sequences are represented with thin lines. Relevant restriction sites sequences are indicated. The position of the *tcf*-operon, i.e. *tcfA*, *B*, *C* and *D* [(SEQ ID NO 2)] (SEQ ID NO:6) is represented by a shaded arrow.

Paragraph at page 4, lines 15-21

Sequence listing

[(SEQ ID NO 1)] SEQ ID NO:1 - DNA sequence of the genes encoding the precursor of the *saf* fimbriae unit of *Salmonella enterica* subspecies I.

[(SEQ ID NO 2)] SEQ ID NO:6 - DNA sequence of the genes which encode the precursor of the *tcf* fimbriae unit of *Salmonella enterica* subspecies I serovar Typhi.

Paragraph at page 4, lines 23-29

Deposit information

The phages carrying the inserted [(SEQ ID NO 1)] SEQ ID NO:1, i.e. phages clones B1, D1, F11 and N10 (see Figure 1) have been given the ECACC Accession numbers 99051922, 99051923, 99051924, and 99051925, respectively.

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The cosmide carrying the inserted [SEQ ID NO 2] SEQ ID NO:6, i.e. cosmide pTY52 (see Figure 2) has been given the ECACC Accession number 99051926.



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Attachment to Supplemental Preliminary Amendment filed July 8, 2002

Mark-up of Claims

1. (Amended) [Peptide] A peptide encoded by a nucleotide sequence selected from [Sequence Listing No. 1 (SEQ ID NO 1)] SEQ ID NO:1 and [Sequence Listing No. 2 (SEQ ID NO 2)] SEQ ID NO:6 for use in medicine.

2. (Amended) Antibodies directed against a peptide encoded by a nucleotide sequence selected from [Sequence Listing No. 1 (SEQ ID NO 1)] SEQ ID NO:1 and [Sequence Listing No. 2 (SEQ ID NO 2)] SEQ ID NO:6 for use in medicine.

3. (Amended) [Nucleotide] A nucleotide sequence selected from [Sequence Listing No. 1 (SEQ ID NO 1)] SEQ ID NO:1 and [Sequence Listing No. 2 (SEQ ID NO 2)] SEQ ID NO:6 for use in medicine.

4. (Amended) A vaccine for the protection against diseases caused by *Salmonella enterica* subspecies I, comprising a peptide encoded by a nucleotide sequence selected from [Sequence Listing No. 1 (SEQ ID NO 1)] SEQ ID NO:1, or antibodies directed against said peptide and, optionally, a pharmaceutically acceptable carrier.

5. (Amended) A vaccine for the protection against diseases caused by *Salmonella enterica* subspecies I serovar Typhi, comprising a peptide encoded by a nucleotide sequence selected from [Sequence Listing No. 2 (SEQ ID NO 2)] SEQ ID NO:6 or antibodies directed against said peptide and, optionally, a pharmaceutically acceptable carrier.

Attachment to Supplemental Preliminary Amendment filed July 8, 2002

6. (Amended) A nucleic acid vaccine for the protection against diseases caused by *Salmonella enterica* subspecies I, comprising a nucleic acid sequence selected from [Sequence Listing No. 1 (SEQ ID NO 1)] SEQ ID NO:1 and, optionally, a pharmaceutically acceptable carrier.

7. (Amended) A nucleic acid vaccine for the protection against diseases caused by *Salmonella enterica* subspecies I serovar Typhi, comprising a nucleic acid sequence selected from [Sequence Listing No. 2 (SEQ ID NO 2)] SEQ ID NO:6 and, optionally, a pharmaceutically acceptable carrier.

8. (Amended) A vector vaccine for the protection against diseases caused by *Salmonella enterica* subspecies I, comprising a host in which a recombinant vector comprising a nucleic acid sequence selected from [Sequence Listing No. 1 (SEQ ID NO 1)] SEQ ID NO:1, has been inserted and, optionally, a pharmaceutically acceptable carrier.

9. (Amended) A vector vaccine for the protection against diseases caused by *Salmonella enterica* subspecies I serovar Typhi, comprising a host in which a recombinant vector comprising a nucleic acid sequence selected from [Sequence Listing No. 2 (SEQ ID NO 2)] SEQ ID NO:6, has been inserted and, optionally, a pharmaceutically acceptable carrier.

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10. A method for vaccinating a mammal against diseases caused by *Salmonella enterica* subspecies I, comprising administering a vaccine according to claim 4 to said mammal.
11. A method for vaccinating a mammal against diseases caused by *Salmonella enterica* subspecies I serovar Typhi, comprising administering a vaccine according to claim 5 to said mammal.
12. (Amended) Antibodies directed against a peptide encoded by a nucleotide sequence selected from [Sequence Listing No. 1 (SEQ ID NO 1)] SEQ ID NO:1, and from [Sequence Listing No. 2 (SEQ ID NO 2)] SEQ ID NO:6, for use in a diagnostic method.
13. (Amended) Peptide encoded by a nucleotide sequence selected from [Sequence Listing No. 1 (SEQ ID NO 1)] SEQ ID NO:1, and [Sequence Listing No. 2 (SEQ ID NO 2)] SEQ ID NO:6, for use in a diagnostic method.
14. (Amended) Primers for, or probes that hybridize with, a nucleotide sequence selected from [Sequence Listing No. 1 (SEQ ID NO 1)] SEQ ID NO:1 and [Sequence Listing No. 2 (SEQ ID NO 2)] SEQ ID NO:6, for use in a diagnostic method for the purpose of detecting *Salmonella enterica* subspecies I.

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15. A method for vaccinating a mammal against diseases caused by *Salmonella enterica* subspecies I, comprising administering a vaccine according to claim 6 to said mammal.

16. A method for vaccinating a mammal against diseases caused by *Salmonella enterica* subspecies I, comprising administering a vaccine according to claim 8 to said mammal.

17. A method for vaccinating a mammal against diseases caused by *Salmonella enterica* subspecies I serovar Typhi, comprising administering a vaccine according to claim 7 to said mammal.

18. A method for vaccinating a mammal against diseases caused by *Salmonella enterica* subspecies I, comprising administering a vaccine according to claim 9 to said mammal.



Patent

Attorney's Docket No. 033267-011

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

In re Patent Application of) BOX: SEQUENCE
)
Anders FOLKESSON et al.)
)
Application Serial No.: 09/926,596) Group Art Unit:
)
Filed: November 26, 2001) Examiner:
)
For: FIMBRIAL PROTEINS) Confirmation No.: 6671

DECLARATION PURSUANT TO
37 C.F.R. §§ 1.821-825

Assistant Commissioner for Patents
Washington, D.C. 20231

Sir:

I, Sharon E. Crane, declare as follows:

1. That the content of the paper copy of the sequence listing and the computer readable copy of the sequence listing submitted in accordance with 37 C.F.R. § 1.821(c) and (e), respectively, are the same in compliance with 37 C.F.R. §§ 1.821(f).
2. That the submission, filed in accordance with 37 C.F.R. §§ 1.821(g)[or (h)] & 1.825(a), herein does not include new matter [or go beyond the disclosure in the international application].

I hereby declare that all statements made herein of my own knowledge are true and that all statements were made on information and belief and are believed to be true; and

Declaration Pursuant to 37 C.F.R. §§ 1.821-.825

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further that these statements were made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under Section 1001 of Title 18 of the United States Code and that such willful false statements may jeopardize the validity of the application or any patent issued.

July 8, 2002

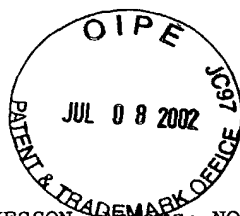
Date



Sharon E. Crane, Ph.D.

Reg. No. 36,113

PTO/PCT Rec'd 08 JUL 2002



SEQUENCE LISTING

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Met Lys Ser
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Ile Lys Lys Leu Ile Ile Ala Ser Ala Leu Ser Met Met Ala Ala Ser
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Cys Tyr Ala Gly Ser Phe Leu Pro Asn Ser Glu Gln Gln Lys Ser Val
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gat att gtg ttt tcc tct ccc caa gat tta acc gta tcg ctt att cca 37520
Asp Ile Val Phe Ser Ser Pro Gln Asp Leu Thr Val Ser Leu Ile Pro
40 45 50
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Val Ser Gly Leu Lys Ala Gly Lys Asn Ala Pro Ser Ala Lys Ile Ala
55 60 65
aag ctt gta gtt aat tct act act ctt aaa gaa ttc ggg gtc agg ggg 37616
Lys Leu Val Val Asn Ser Thr Thr Leu Lys Glu Phe Gly Val Arg Gly
70 75 80
att tct aac aac gtg gta gac agt act ggc act gca tgg cgt gta gct 37664
Ile Ser Asn Asn Val Val Asp Ser Thr Gly Thr Ala Trp Arg Val Ala
85 90 95
ggg aaa aat act ggt aaa gag atc ggt gtg ggc tta tca agt gac agt 37712
Gly Lys Asn Thr Gly Lys Glu Ile Gly Val Gly Leu Ser Ser Asp Ser
100 105 110 115

$$\frac{1}{\Gamma(\alpha)} \int_0^t (t-\tau)^{\alpha-1} f(\tau) d\tau = I^\alpha f(t), \quad t > 0, \quad \alpha > 0$$

gat gtt gca gga aat tta aga tgg gtg gag acg ggc aac aaa ctt aag	38491
Asp Val Ala Gly Asn Leu Arg Trp Val Glu Thr Gly Asn Lys Leu Lys	
335 340 345	
gtg gag aac ccc acc ccg ttt tac atg aat tta gcc tct gtc aca gta	38539
Val Glu Asn Pro Thr Pro Phe Tyr Met Asn Leu Ala Ser Val Thr Val	
350 355 360	
ggg gga aag ccc att aca ggg ctt gag tat gtc ccc ccc ttt gct gac	38587
Gly Gly Lys Pro Ile Thr Gly Leu Glu Tyr Val Pro Phe Ala Asp	
365 370 375	
aaa aca cta aat atg cca ggt agt gcc cat ggt gat atc gag tgg aga	38635
Lys Thr Leu Asn Met Pro Gly Ser Ala His Gly Asp Ile Glu Trp Arg	
380 385 390 395	
gtt att aca gac ttt ggt ggt gaa agt cat ccg ttc cac tac gtt ctt	38683
Val Ile Thr Asp Phe Gly Gly Glu Ser His Pro Phe His Tyr Val Leu	
400 405 410	
aaa taa atccaggggc ttagcggcag aaa atg aag ttc aaa caa cct gcc ttg	38736
Lys Met Lys Phe Lys Gln Pro Ala Leu	
415 420	
cta ctg ttc atc gcg gga gtg gtt cat tgc gca aat gcg cac act tac	38784
Leu Leu Phe Ile Ala Gly Val Val His Cys Ala Asn Ala His Thr Tyr	
425 430 435	
aca ttc gat gca tca atg ttg ggc gat gca gcg aaa ggg gtt gat atg	38832
Thr Phe Asp Ala Ser Met Leu Gly Asp Ala Ala Lys Gly Val Asp Met	
440 445 450	
tcg ctc ttt aac cag ggg tta caa cag cca ggg act tat cgc gtg gac	38880
Ser Leu Phe Asn Gln Gly Leu Gln Gln Pro Gly Thr Tyr Arg Val Asp	
455 460 465	
gtg atg gtg aat ggt aaa cgt gtc gac acc cgt gat gtg gtg ttc aaa	38928
Val Met Val Asn Gly Lys Arg Val Asp Thr Arg Asp Val Val Phe Lys	
470 475 480 485	
ttg gaa aag gat ggg caa gga acg cct gtt ctg gct cct tgt ttg acg	38976
Leu Glu Lys Asp Gly Gln Gly Thr Pro Val Leu Ala Pro Cys Leu Thr	
490 495 500	
gtc agt cag ctt tca cgc tac ggc gta aaa acg gaa gat tac cct cag	39024
Val Ser Gln Leu Ser Arg Tyr Gly Val Lys Thr Glu Asp Tyr Pro Gln	
505 510 515	
ttg tgg aaa gca gca aag ccc cca gat gag tgt gcg gat ctg acc gcc	39072
Leu Trp Lys Ala Ala Lys Pro Pro Asp Glu Cys Ala Asp Leu Thr Ala	
520 525 530	
att cca cag gct aaa gcg gta ctg gat atc aat aat cag caa ctg caa	39120
Ile Pro Gln Ala Lys Ala Val Leu Asp Ile Asn Asn Gln Gln Leu Gln	
535 540 545	
ctg agt att ccg cag ttg gcg ttg cgt ccg gaa ttt aag ggg atc gct	39168
Leu Ser Ile Pro Gln Leu Ala Leu Arg Pro Glu Phe Lys Gly Ile Ala	
550 555 560 565	

cca gaa gat ctt tgg gat gat ggt att ccg gcg ttt ctg atg aac tac	39216
Pro Glu Asp Leu Trp Asp Asp Gly Ile Pro Ala Phe Leu Met Asn Tyr	
570 575 580	
agt gcg agg aca acg cag acg gat tac aaa atg gat atg gtg ggg cgt	39264
Ser Ala Arg Thr Thr Gln Thr Asp Tyr Lys Met Asp Met Val Gly Arg	
585 590 595	
gac aac tct tcc tgg gta caa ctg caa ccg gga atc aat ata ggt gcg	39312
Asp Asn Ser Ser Trp Val Gln Leu Gln Pro Gly Ile Asn Ile Gly Ala	
600 605 610	
tgg cgt gtc cgc aat gcg acc agc tgg cag ccg agt agt caa ctg tcg	39360
Trp Arg Val Arg Asn Ala Thr Ser Trp Gln Arg Ser Ser Gln Leu Ser	
615 620 625	
ggg aag tgg cag gca gca tat acc tat gct gag cgt gga ctg tac tca	39408
Gly Lys Trp Gln Ala Ala Tyr Thr Tyr Ala Glu Arg Gly Leu Tyr Ser	
630 635 640 645	
cta aaa agt cgt ctg act ctg ggg caa aag act tcg cag ggg gag ata	39456
Leu Lys Ser Arg Leu Thr Leu Gly Gln Lys Thr Ser Gln Gly Glu Ile	
650 655 660	
ttt gat agt gtg cca ttt acc ggt gtg atg ttg gca tcg gat gac aac	39504
Phe Asp Ser Val Pro Phe Thr Gly Val Met Leu Ala Ser Asp Asp Asn	
665 670 675	
atg gtg ccc tac agt gag ccg cag ttt gct ccg gta gtg cgt ggg att	39552
Met Val Pro Tyr Ser Glu Arg Gln Phe Ala Pro Val Val Arg Gly Ile	
680 685 690	
gcc cgc acg cag gct ccg gtg gag gtc aaa cag aat ggt tac acc att	39600
Ala Arg Thr Gln Ala Arg Val Glu Val Lys Gln Asn Gly Tyr Thr Ile	
695 700 705	
tac aac acc act gtg gcg ccc gga ccg ttt gca ctg ccg gat ctg tcg	39648
Tyr Asn Thr Thr Val Ala Pro Gly Pro Phe Ala Leu Arg Asp Leu Ser	
710 715 720 725	
gta aca gac agt agt ggt gat ctg cat gtc acc gtg tgg gag gcc gat	39696
Val Thr Asp Ser Ser Gly Asp Leu His Val Thr Val Trp Glu Ala Asp	
730 735 740	
ggc agt aca caa atg ttt gtg gtg ccg tat cag acc ccg gcg ata gca	39744
Gly Ser Thr Gln Met Phe Val Val Pro Tyr Gln Thr Pro Ala Ile Ala	
745 750 755	
ctg cac cag gga tat ttg aag tac agc ctg ttg gcg ggc cga tac cga	39792
Leu His Gln Gly Tyr Leu Lys Tyr Ser Leu Leu Ala Gly Arg Tyr Arg	
760 765 770	
tcg tca gac tct gca acg gat aag ccg cag atc gcg cag gct acg ttg	39840
Ser Ser Asp Ser Ala Thr Asp Lys Arg Gln Ile Ala Gln Ala Thr Leu	
775 780 785	
atg tat ggt ctg ccg tgg aat ctc act gca tac ggc ggt ata cag agt	39888
Met Tyr Gly Leu Pro Trp Asn Leu Thr Ala Tyr Gly Gly Ile Gln Ser	
790 795 800 805	

gca acg cat aat caa gct gca ttg ctt ggt ttg ggg gga tct ctc ggg	39936
Ala Thr His Asn Gln Ala Ala Leu Leu Gly Leu Gly Gly Ser Leu Gly	
810 815 820	
cgg tgg ggg agt tta tct gtc gat gga agc gac aca cac agt cag cgt	39984
Arg Trp Gly Ser Leu Ser Val Asp Gly Ser Asp Thr His Ser Gln Arg	
825 830 835	
cag ggg gag gcg gta cag caa gga gcc tcc tgg cga ctg cgt tac agc	40032
Gln Gly Glu Ala Val Gln Gln Gly Ala Ser Trp Arg Leu Arg Tyr Ser	
840 845 850	
aac cag ctg act gcg acg ggg aca aat ttt ttt ctg acg aga tgg cag	40080
Asn Gln Leu Thr Ala Thr Gly Thr Asn Phe Phe Leu Thr Arg Trp Gln	
855 860 865	
tat gcc tcg cag ggc tat aac acc cta tcc gat gtg ctc gac agt tat	40128
Tyr Ala Ser Gln Gly Tyr Asn Thr Leu Ser Asp Val Leu Asp Ser Tyr	
870 875 880 885	
cga cat aat ggc aac cgt cta tgg tcg tgg cgg gaa aat ttg cag ccg	40176
Arg His Asn Gly Asn Arg Leu Trp Ser Trp Arg Glu Asn Leu Gln Pro	
890 895 900	
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Ser Ser Arg Thr Thr Leu Met Leu Ser Gln Ser Trp Gly Arg His Leu	
905 910 915	
ggc aat ctg agt tta acc ggt tcc cgt acc gac tgg cgt aat cgc ccc	40272
Gly Asn Leu Ser Leu Thr Gly Ser Arg Thr Asp Trp Arg Asn Arg Pro	
920 925 930	
ggt cat gat gac agc tac gga ctg agt tgg gga acc tct atc gga ggg	40320
Gly His Asp Asp Ser Tyr Gly Leu Ser Trp Gly Thr Ser Ile Gly Gly	
935 940 945	
ggc tcg ctg tca ttg aac tgg aat caa aac aga acg ctg tgg cgc aat	40368
Gly Ser Leu Ser Leu Asn Trp Asn Gln Asn Arg Thr Leu Trp Arg Asn	
950 955 960 965	
ggc gcg cac cgt aaa gag aac ata acc agc ctg tgg ttc agt atg cca	40416
Gly Ala His Arg Lys Glu Asn Ile Thr Ser Leu Trp Phe Ser Met Pro	
970 975 980	
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Leu Ser Arg Trp Thr Gly Asn Asn Val Ser Ala Ser Trp Gln Met Thr	
985 990 995	
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Ser Pro Ser His Gly Gly Gln Thr Gln Gln Val Gly Val Asn Gly Glu	
1000 1005 1010	
gca ttc agt cag caa ctg gat tgg gag gtg cgt cag agt tac cgt gcc	40560
Ala Phe Ser Gln Gln Leu Asp Trp Glu Val Arg Gln Ser Tyr Arg Ala	
1015 1020 1025	
gat gcc ccg cca ggt ggt ggt aat aac agc gca ttg cac ttg gca tgg	40608
Asp Ala Pro Pro Gly Gly Gly Asn Asn Ser Ala Leu His Leu Ala Trp	
1030 1035 1040 1045	

aat ggg gat tac ggc ctg tta ggt ggt gac tat agc tac agc cgg gcg	40656
Asn Gly Asp Tyr Gly Leu Leu Gly Gly Asp Tyr Ser Tyr Ser Arg Ala	
1050 1055 1060	
atg cgc cag atg gga gtc aat atc gcg gga ggt ata gtt atc cac cat	40704
Met Arg Gln Met Gly Val Asn Ile Ala Gly Gly Ile Val Ile His His	
1065 1070 1075	
cat ggt gtg acg ctg ggg caa cct ttg caa ggc tca gtg gcg ctg gtt	40752
His Gly Val Thr Leu Gly Gln Pro Leu Gln Gly Ser Val Ala Leu Val	
1080 1085 1090	
gaa gcg cca ggg gcc tcg ggg gtg cca gtt ggc ggc tgg cct ggc gtt	40800
Glu Ala Pro Gly Ala Ser Gly Val Pro Val Gly Gly Trp Pro Gly Val	
1095 1100 1105	
aag acg gat ttt cgt ggc gac acc aca gtg ggc aac ctg aac gtc tat	40848
Lys Thr Asp Phe Arg Gly Asp Thr Thr Val Gly Asn Leu Asn Val Tyr	
1110 1115 1120 1125	
cag gag aat aca gtc agc ctc gat ccg tcg cga cta ccg gat gac gca	40896
Gln Glu Asn Thr Val Ser Leu Asp Pro Ser Arg Leu Pro Asp Asp Ala	
1130 1135 1140	
gag gtc aca caa acc gat gtg cgc gtg gtg cca acc gaa ggg gcg gtg	40944
Glu Val Thr Gln Thr Asp Val Arg Val Val Pro Thr Glu Gly Ala Val	
1145 1150 1155	
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Val Glu Ala Lys Phe His Thr Arg Ile Gly Ala Arg Ala Leu Met Thr	
1160 1165 1170	
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Leu Lys Arg Glu Asp Gly Ser Ala Ile Pro Phe Gly Ala Gln Val Thr	
1175 1180 1185	
gtc aat ggg cag gat ggc agt gct gct ctg gtg gat act gat agc cag	41088
Val Asn Gly Gln Asp Gly Ser Ala Ala Leu Val Asp Thr Asp Ser Gln	
1190 1195 1200 1205	
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Val Tyr Leu Thr Gly Leu Ala Asp Lys Gly Glu Leu Thr Val Lys Trp	
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Gly Ala Gln Gln Cys Arg Val Asn Tyr Arg Leu Pro Ala His Lys Gly	
1225 1230 1235	
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Ile Ala Gly Leu Tyr Gln Met Ser Gly Leu Cys Arg	
1240 1245 1250	
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Met Trp Met Lys Ile Gln Arg Val Lys Thr Val Ile Tyr	
1255 1260	
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Ser Val Ser Leu Leu Val Ala Ala Ser Ser Leu Val Pro Ile Ala Asn	
1265 1270 1275	

gcc gca gaa aaa ctt cag aca acg cta cgt gta ggt act tac ttt cgg 41379
Ala Ala Glu Lys Leu Gln Thr Thr Leu Arg Val Gly Thr Tyr Phe Arg
1280 1285 1290 1295

gct ggg cac gtg cca gat ggg atg gtg ctt gcg caa ggc tgg gtg act 41427
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Date of Deposit: 26 November 2001

I hereby certify that this paper or fee is being hand-deposited with the United States Patent and Trademark Office on the date indicated above.

Jennifer Harris-Lohse
Typed or Printed Name of Person Depositing Paper

Jennifer Harris-Lohse
Signature of Person Depositing Paper

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

In Re Application of:

FOLKESSON et al.

Serial No.: PCT/SE00/01079

Art Unit: (unassigned)

Filing Date: Herewith

Examiner: (unassigned)

Title: FIMBRIAL PROTEINS

PRELIMINARY AMENDMENT

Commissioner for Patents
Washington, D.C. 20231

Sir:

Prior to calculating the filing fee and examination of the above-identified application on the merits, please enter the following amendments.

AMENDMENT

In the Claims:

Please amend claims 10 and 11 as follows.

10. (Amended) A method for vaccinating a mammal against diseases caused by *Salmonella enterica* subspecies I, comprising administering a vaccine according to claim 4 to said mammal.

11. (Amended) A method for vaccinating a mammal against diseases caused by *Salmonella enterica* subspecies I serovar Typhi, comprising administering a vaccine according to claim 5 to said mammal.

Please add the following new claims.

15. (New) A method for vaccinating a mammal against diseases caused by *Salmonella enterica* subspecies I, comprising administering a vaccine according to claim 6 to said mammal.

16. (New) A method for vaccinating a mammal against diseases caused by *Salmonella enterica* subspecies I, comprising administering a vaccine according to claim 8 to said mammal.

17. (New) A method for vaccinating a mammal against diseases caused by *Salmonella enterica* subspecies I serovar Typhi, comprising administering a vaccine according to claim 7 to said mammal.

18. (New) A method for vaccinating a mammal against diseases caused by *Salmonella enterica* subspecies I serovar Typhi, comprising administering a vaccine according to claim 9 to said mammal.

VERSION WITH MARKINGS TO SHOW CHANGES MADE

In the Claims:

Claims 10 and 11 have been amended as follows.

10. (Amended) A method for [protection] vaccinating a mammal against diseases caused by *Salmonella enterica* subspecies I, comprising administering a vaccine according to [any of claims] claim 4[, 6, and 8] to said mammal.

11. (Amended) A method for [protection] vaccinating a mammal against diseases caused by *Salmonella enterica* subspecies I serovar Typhi, comprising administering a vaccine according to [any of claims] claim 5[, 7, and 9] to said mammal.

New claims 15-18 have been added.

--15. (New) A method for vaccinating a mammal against diseases caused by *Salmonella enterica* subspecies I, comprising administering a vaccine according to claim 6 to said mammal.

16. (New) A method for vaccinating a mammal against diseases caused by *Salmonella enterica* subspecies I, comprising administering a vaccine according to claim 8 to said mammal.

17. (New) A method for vaccinating a mammal against diseases caused by *Salmonella enterica* subspecies I serovar Typhi, comprising administering a vaccine according to claim 7 to said mammal.

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PATENT

18. (New) A method for vaccinating a mammal against diseases caused by *Salmonella enterica* subspecies I serovar Typhi, comprising administering a vaccine according to claim 9 to said mammal.--

REMARKSIntroductory Comments:

The present application is a 35 U.S.C. §371 filing of International Application Number PCT/SE00/01079, filed 26 May 2000.

Overview of the Amendments:

Applicants, by way of this preliminary amendment, have entered minor claim amendments and four new claims for consideration herein. Applicants respectfully request that these amendments be entered prior to calculation of the filing fees, and prior to examination of the present application on the merits.

More particularly, applicants have amended claims 10 and 11 merely to remove multiple dependencies and to recite the invention with greater particularity. Support for these claim amendments can be found in the claims as originally filed and throughout the specification, for example, at page 6, line 15 through page 7, line 11. In addition, applicants have submitted herewith new claims 15-18. Support for these new claims can be found throughout the specification and claims, particularly in claims 10 and 11 as originally filed and in the specification at page 6, line 15 through page 7, line 11. Accordingly, applicants submit that no new matter has been added by way of the above-described claim amendments and the new claims, and the entry thereof is respectfully requested.

The claim amendments and the new claims have been provided in both "clean version" and in "marked-up version" in conformance with 37 C.F.R. §1.121(c). The "marked-up version" of these amendments and new claims shows the changes made to the claims by the current amendment. The attached pages are captioned "VERSION WITH MARKINGS TO SHOW CHANGES MADE".

Applicants respectfully submit that the claims define an invention which complies with the requirements of 35 U.S.C. § 112 and which is novel and nonobvious over the art. Accordingly, allowance is believed to be in order and an early notification to that effect would be appreciated.

By: Thomas P. McCracken
Thomas P. McCracken
Registration No. 38,548

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FIMBRIAL PROTEINSField of the invention

The present invention is based on the finding that two fimbrial operons, the *saf* operon and the *tcf* operon, are specific for *Salmonella enterica* subspecies 1 bacteria and therefore have therapeutic use. Due to their specificity they can be used to provide vaccines against *Salmonella enterica* subspecies I as well as for detection of *Salmonella enterica* subspecies I. The *saf* operon is specific for all *Salmonella enterica* subspecies 1 bacteria and the *tcf* operon is specific for the serovar Typhi of *Salmonella enterica* subspecies 1.

All or part of the DNA-sequences of the genes encoding these proteins can be used as active agents in a vaccine against diseases caused by the *Salmonella enterica* subspecies I bacterial strains or for detection of said bacterial strains.

The present invention also relates to methods of isolating these fimbrial proteins, to antibodies directed against these proteins, and to a vaccine composition comprising these proteins or antibodies directed against these proteins for use in the treatment of infections caused by the *Salmonella spp.* The fimbrial proteins according to the invention or antibodies directed against them can be used for detection of *Salmonella spp.* bacteria.

Background of the invention

The members of genus *Salmonella spp* colonize and infect a wide range of different organisms. Many cause gastroenteritis and enteric fever in humans and domesticated animals while others are not associated with human disease (Saylers et al, 1994). The genus has been divided into two species, *Salmonella bongori* and *Salmonella enterica* where *enterica* can be further subdivided into seven subspecies, designated I, II, IIIa, IIIb, IV, VI, and VII (Reeves et al, 1989). *Salmonella enterica* subspecies I are preferentially associated with warm-blooded animals. Over 99% of all clinical *Salmonella* isolates are strains belonging to this subspecies, including serovars Typhimurium and Enteritidis, which are the major causes of *Salmonella* induced gastroenteritis in humans, and Typhi, the human specific causative organism of typhoid fever, the most severe form of human salmonellosis (Popoff et al, 1992).

Salmonella enterica subspecies I consists of over 1300 different serovars and is preferentially associated with warm-blooded animals (Bäumler, 1997). Over 99% of all clinical *Salmonella* isolates are strains belonging to this subspecies,

including serovars Typhimurium and Enteritidis, which are the major causes of *Salmonella* induced gastroenteritis in humans, and Typhi, the human specific causative organism of typhoid fever, the most severe form of human salmonellosis (Popoff and Le Minor, 1992).

5

Today gastroenteritis and enteric fever can neither be prevented nor treated with good results. Typhoid fever is a substantial public health problem in developing countries. Each year 33 million people become ill and over 500 000 people die from this infection (American Institute of Medicine, 1986). Typhoid fever can be prevented by vaccination with attenuated bacteria, such as Ty21 and Vi vaccines and whole cell vaccines. Whole cell vaccines show a high incidence of side effects (Ashcroft et al, 1964, Yugoslav Typhoid commission, 1964). The vaccines consisting of attenuated strains of *Salmonella typhi* suffer from serious drawbacks. They must be administered as three or four spaced doses in order to stimulate protective immune responses (Levine et al, 1989). The treatment of *Salmonella typhi* with antibiotics is jeopardized since there are strains of *Salmonella typhi* that are resistant to chloramphenicol, ampicillin, and trimethoprim as well as ciprofloxacin (i.e. multidrug-resistant strains) (Rowe et al, 1997).

20

Accurate detection of *Salmonella enterica* subspecies I is today not possible. *Salmonella enterica* subspecies I can today only be detected by antibodies directed against surface proteins of *Salmonella enterica* subspecies I. The use of the sequences according to the invention makes it for the first time possible to rapidly and accurately determine the presence of *Salmonella enterica* subspecies I.

25

For many pathogenic bacteria, there is evidence that the filamentous surface protein structures called pili (fimbriae) are connected to the adhesion of the bacteria to the host cells. Pili proteins are very antigenic and are easily purified. Therefore pili preparations have been used as antigens for vaccination.

30

Summary of the invention

35

The invention relates to the objects as defined in the claims. The main object of the present invention is to provide two fimbrial proteins that are specific for *Salmonella enterica* subspecies I bacterial strains, the nucleotide sequences

encoding said proteins, as well as the corresponding amino acid sequences of for therapeutic and diagnostic use. Further are recombinant microorganisms provided, in which the nucleotide sequences according to the invention have been inserted.

5 An object of the present invention is to provide vaccine compositions for use in the treatment of *Salmonella enterica* infective strains, essentially pure Saf and Tcf fili protein of *Salmonella enterica* subspecies I and *Salmonella enterica* subspecies I serovar Typhi, respectively, as well as antibodies directed to these
10 fili proteins.

A further object of the present invention is to provide the DNA sequences of the genes encoding the Saf and Tcf proteins. These sequences can be used for recombinant production of the proteins and for the preparations of vector
15 vaccines against *Salmonella enterica* subspecies 1 and *Salmonella enterica* subspecies 1 serovar Typhi, respectively, as well as for diagnostic purposes.

Yet another object of the present invention to use purified Saf and Tcf protein from *Salmonella enterica* subspecies 1 bacteria for active or passive
20 immunization of mammals, i.e. the proteins according to the invention can be comprised in a vaccine composition or be used to raise antibodies which can be comprised in a vaccine composition.

Finally, an object of the present invention is to provide a method for preventing
25 or reducing the possibility of *Salmonella* infection of a mammal by administering the vaccines according to the invention. The invention may be more fully understood by reference to the following drawings and detailed description.

30 Brief description of the drawings

Figure 1.

Schematic representation of phage clones (named N10, D1, B1, F11) covering the entire cs7 insert of *Salmonella enterica* serovar Typhimurium strain SR χ 3181, i.e. comprising the *saf* fimbrial operon, i.e. *safA*, *B*, *C* and *D*
35 (SEQ ID NO 1).

The clones were selected from partial *Eco* RI and *Bam* HI libraries in the Lambda Dash II vector. The cs7 insert is represented by a bold line. The extent

of respective phage insert is represented by horizontal bars. Name and size of the phage inserts are indicated on the left side of the figure.

Figure 2: Schematic representation of the pTY52 cosmid comprising the *tcf*-operon (SEQ ID NO 2).

- 5 A *tcf* specific PCR fragment of 11105 bp was cloned into the Expand vector I cosmid (Roche). The insert is represented with a thick black line while vector sequences are represented with thin lines. Relevant restriction sites sequences are indicated. The position of the *tcf*-operon, i.e. *tcfA*, *B*, *C* and *D* (SEQ ID NO 2), is represented by a shaded arrow.
- 10 Figure 3: The phylogenetic distribution of the identified genes on the cs7 insert was investigated using the well defined SARC collection, see Example 1.
- Figure 4: A 2 kb large internal *EcoR* I fragment was used as a probe in a Southern blot of the SARC collection, see Example 2.

15 Sequence listing

Sequence Listing No. 1, herein referred to as SEQ ID NO 1, —DNA sequence of the genes encoding the precursor of the *saf* fimbriae unit of *Salmonella enterica* subspecies I.

- Sequence Listing No. 2, herein referred to as SEQ ID NO 2, —DNA sequence of
- 20 the genes which encode the precursor of the *tcf* fimbriae unit of *Salmonella enterica* subspecies I serovar Typhi.

Deposit information

- The phages carrying the inserted SEQ ID NO 1, i.e. phages clones B1, D1, F11
- 25 and N10 (see Figure 1) have been given the ECACC Accession numbers 99051922, 99051923, 99051924, and 99051925, respectively.

The cosmid carrying the inserted SEQ ID NO 2, i.e. cosmid pTY52 (see Figure 2) has been given the ECACC Accession number 99051926.

The depositions were made May 19, 1999.

30

Detailed description of the invention

- The present invention is based on the finding that two fimbrial operons, the *saf* operon and the *tcf* operon, are specific for *Salmonella enterica* subspecies I bacteria. Due to their specificity they can be used to provide vaccines against
- 35 *Salmonella enterica* subspecies I as well as detection methods for *Salmonella enterica* subspecies I. The *saf* operon is specific for all *Salmonella enterica*

subspecies 1 bacteria and the *tcf* operon is specific for the serovar typhi of *Salmonella enterica* subspecies 1, see Examples 1 & 2.

The main object of the invention relates to two fimbrial operons, the *saf* operon
5 and the *tcf* operon, that are specific for *Salmonella enterica* subspecies 1 bacteria for therapeutic use.

Another object of the present invention is to provide vaccines against *Salmonella enterica* subspecies 1 induced gastroenteritis, enteric fever and
10 typhoid fever.

A further object of the present invention is to provide methods to detect *Salmonella enterica* subspecies 1. The nucleotide sequences according to the invention are useful for constructing vectors for use as vaccines for insertion
15 into attenuated bacteria in constructing a recombinant vaccine, for insertion into a viral vector in constructing a recombinant viral vaccine, or for direct inoculation as a nucleic acid vaccine. The pili proteins according to the invention, or antigenic fragments thereof, can be used for active immunization and antibodies directed against them can be used for passive immunization. All
20 these applications of the sequences according to the invention are obtained by applying standard techniques known to the man ordinary skilled in the art.

Vaccines against *Salmonella enterica* subspecies I.

The genes encoding the *saf* and *tcf* fimbrial structures, or fragments thereof,
25 may be incorporated into a bacterial or viral vaccine comprising recombinant bacteria, virus or fungi which are engineered to produce one or more immunogenic epitopes of the *saf* or *tcf* fimbrial structures. In addition, the genes encoding the *saf* and *tcf* fimbrial structures, or part thereof, operatively linked to regulatory elements, can be introduced directly as a nucleic acid
30 vaccine, to elicit a protective immune response.

The proteins or antigenic fragment thereof, deduced from the nucleic acid sequences of the present invention are useful alone or in conventional vaccine mixtures in the vaccine compositions according to the invention. The proteins
35 could be produced by chemical synthesis or recombinant expression according to conventional methods.

The different vaccines according to the present invention are administered to mammals in many different ways. These include intradermal, intramuscular, intraperitoneal, intravenous, subcutaneous, oral, and intranasal routes of administration. The vaccine doses will differ depending on circumstances such

as body weight, interferences with other administered medicaments etc. The upper limit is not critical unless the dose shows toxicity.

The peptides and proteins of the present invention are also useful to produce
5 monoclonal or polyclonal antibodies for use in passive immunotherapy against *Salmonella enterica* subspecies 1. Human immunoglobulin is preferred. Antisera is obtained from individuals immunized with proteins or peptides according to the invention. The immunoglobulin fraction is then enriched, for example by immunoaffinity or affinity chromatography. Antibodies raised in
10 a suitable mammal or in the patient to be treated, can subsequently be administered locally or topically, e.g. orally to the patient.

Detection of *Salmonella enterica* subspecies I in general.

The sequences according to the invention, or part thereof, or fragments
15 hybridizing therewith, as well as the proteins according to the invention, or part thereof, and antibodies directed to said proteins, or antigenic fragments thereof, can be used in molecular diagnostic assays for the detection of *Salmonella enterica* subspecies I.

20 Nucleic acids having the nucleotide sequence according to the invention, or any nucleotide sequence hybridizing therewith can be used as a probe in nucleic acid hybridization assays for the detection of *Salmonella spp* in various tissues and body fluids of patients. The hybridization assay may be of any type including; Southern blots, Northern blots, colony blots.

25 PCR technology is the most preferred technology for detection according to the invention of *Salmonella enterica* subspecies 1. Primers of at least one selected from the 5' end and one from the 3' end can be used in PCR and other known tests to rapidly identify the presence of *Salmonella enterica* subspecies 1. This is
30 according to conventional techniques.

The isolated and purified proteins and peptides of the invention can be used as diagnostics to measure an increase in serum titer of *Salmonella enterica*
35 subspecies I-specific antibody since they bind strongly to these antibodies. A serum test sample can be screened for *Salmonella enterica* subspecies I by methods such as for example ELISA.

The invention further comprises the use of antibodies directed against the *saf* and *tcf* fimbriae structures for quantitative or qualitative determinations of the pili proteins of the invention, or fractions thereof, in cells, tissues or body fluids.

5

Detection of *Salmonella enterica* subspecies I by using nucleic acid hybridization technology

Nucleic acid hybridization technology can also be used to detect *Salmonella enterica* subspecies I according to the invention. The nucleic acid probes chosen from parts of the sequences according to the invention can be either DNA or RNA. DNA sequences complementary to the sequences according to the invention can also be used. The binding of the probe to the target sequence, i.e. the hybridization, must not be perfect. Variations and mutations of the sequences according to the invention can be used as long as they hybridize good enough to detect *Salmonella enterica* subspecies I. The preferred length of the nucleic acid probes is about 10 to 400 nucleotides, most preferred not longer than 100 nucleotides.

The nucleotide probe is preferably chosen from the parts of the sequences that have the least variation. In the most preferred embodiments when screening for SEQ ID NO 1 (the *saf* operon, specific for *Salmonella enterica* subspecies I) a nucleotide probe or PCR primer selected from nucleotides 37 368-37 868 should be avoided since this region is hypervariable.

25 The nucleic acid probes according to the invention are prepared by any conventional method such as organic synthesis, recombinant techniques, or isolation from genomic DNA.

30 The nucleic acid probes of the invention are labeled in a conventional manner to signal hybridization to target nucleic acid from *Salmonella enterica* subspecies I. The labeling may comprise a radiolabel, an enzyme, a bacterial label, a fluorescent label, an antibody, an antigen, a latex particle, an electron dense compound, or a light scattering particle.

35 The probes may be provided in a lyophilized form, to be reconstituted in a buffer appropriate for hybridization, or the probes may already be present in

such a buffer. The buffer may contain a suitable hybridization enhancer, detergent, carrier DNA, and a compound to increase the specificity.

Any conventional hybridization assay technique, such as dot blot hybridization, Southern blotting, sandwich hybridization, displacement hybridization and the like, can be used.

The target analyte polynucleotide of a microorganism may be in various media, most often in a biological, or physiological specimen. In most cases it is preferred to subject the specimen containing the target polynucleotide to any conventional extraction, purification, and/or isolation before conducting the analysis.

The sample containing the target analyte nucleotide sequence must often be treated to convert the DNA to a single-stranded form, which may be accomplished by a variety of conventional techniques, such as thermal or chemical techniques.

The following examples describe the isolation and specificity of the sequences according to the invention.

EXAMPLE 1

Identification and characterization of the *saf* operon.

The present inventors found, upon investigation of a 7 kb chromosomal region on centisome 7 originally isolated from the *S. typhimurium* strain SR-11 k3181, a region that exhibits many of the traits that define a pathogenicity island. It has a lower G+C composition than the average composition of the *Salmonella* genome and includes many sequences related to different mobile genetic elements. The region is not present in *E.coli* K12, and the *Salmonella* specific DNA is inserted between the tRNA gene *aspV* and the stop codon of *yafV*, a hypothetical protein upstream of the *yafH* gene at 5 min in the *E.coli* chromosome. This *Salmonella* specific insert encodes proteins creating adhesive structures and other virulence factors. Sequencing revealed genes encoding a new fimbrial operon that they designated *Salmonella* Atypical Fimbriae (*saf*), due to its relatedness to a subgroup of adhesive structures forming thin atypical fimbriae or non-fimbrial adhesins.

The *saf* operon consists of four contiguous genes, *safA*, *safB*, *safC* and *safD* that encode fimbrial subunit, periplasmic chaperone, outer membrane usher protein and alternative fimbrial subunit, respectively. The genes *safA*, *B*, *C* and *safD* encode putative proteins of 166, 244, 836 and 156 amino acids, respectively. Analyzes of clinical *Salmonella* isolates showed that DNA of 195 out of 198 clinical isolates belonging to *S. enterica* subspecies I hybridized with *safB* and *safC*, i.e. these sequences are common to more than 99% of the known *Salmonella enterica* subspecies 1 bacteria. The inventors showed that 58% of these clinical isolates carry the *safA*, see Table 1.

Table 1. The prevalence of the *saf* genes in clinical *Salmonella* isolates.

Serovar	<i>safA</i>	<i>safB</i>	<i>safC</i>	# isolates
<i>S. adelaide</i>	-	+	+	1
<i>S. agona</i>	+	+	+	6
<i>S. anatum</i>	-	+	+	3
<i>S. bareilly</i>	+	+	+	3
<i>S. blockley</i>	+	+	+	3
<i>S. bovis/morbificans</i>	-	+	+	5
<i>S. braenderup</i>	-	+	+	4
<i>S. brandenburg</i>	+	+	+	1
<i>S. bredeney</i>	+/-	+	+	15
<i>S. chester</i>	+	+	+	1
<i>S. colindale</i>	-	+	+	1
<i>S. derby</i>	-	+	+	1
<i>S. dublin</i>	-	+	+	1
<i>S. eastbourne</i>	+	+	+	2
<i>S. emek</i>	+	+	+	1
<i>S. enteritidis</i>	-	+	+	8
<i>S. give</i>	-	+	+	1
<i>S. goettingen</i>	+	+	+	1
<i>S. haardt</i>	-	+	+	1
<i>S. hadar</i>	+	+	+	16
<i>S. heidelberg</i>	-	+	+	1
<i>S. hittingfoss</i>	+	+	+	5
<i>S. infantis</i>	-/+	+	+	6
<i>S. java</i>	-	+	+	1
<i>S. javiana</i>	-	+	+	1
<i>S. kottbus</i>	-	+	+	1
<i>S. livingstone</i>	-	+	+	1
<i>S. london</i>	+	+	+	1
<i>S. maastricht</i>	+	+	+	2
<i>S. mbandaka</i>	-	-	-	3
<i>S. montevideo</i>	+	+	+	1
<i>S. muenster</i>	-	+	+	1
<i>S. newport</i>	+	+	+	2
<i>S. ohio</i>	+	+	+	1
<i>S. oranienburg</i>	+	+	+	2
<i>S. panama</i>	+	+	+	3
<i>S. potsdam</i>	+	+	+	1
<i>S. rissen</i>	-	-	-	1
<i>S. saarbrücken</i>	-	+	+	1
<i>S. saint paul</i>	+	+	+	3
<i>S. schwartzengrund</i>	-	+	+	1
<i>S. singapore</i>	+	+	+	1
<i>S. stanley</i>	+	+	+	5
<i>S. subsp I 4,5,12:-:-</i>	+	+	+	2
<i>S. subsp I 4,5,12:b:-</i>	-	+	+	1
<i>S. subsp I 4,5,12:i:-</i>	+	+	+	1
<i>S. subsp I sport</i>	-	+	+	1
<i>S. tennessee</i>	+	+	+	2
<i>S. thompson</i>	-	+	+	1
<i>S. typhi</i>	-	+	+	1
<i>S. typhimurium</i>	+	+	+	27
<i>S. virchow</i>	+	+	+	7
<i>S. weltevreden</i>	-	+	+	1
<i>S. worthington</i>	-	-	-	2
<i>S. subsp III</i>	-	-	-	1

The phylogenetic distribution of the identified genes on the *cs7* insert was investigated using the well defined SARC collection, which showed that the presence of the *safA*, *safB*, *safC* and *safD* genes is restricted to *S. enterica* subspecies I (Fig. 3). This region is hence the first subspecies I specific genetic region to be identified with a broad distribution within the subspecies. Since the serovars of subspecies I constitute over 99% of human salmonellosis and are preferentially associated with warm blooded animals, it implicates a role for the *saf* adhesive organelle in the colonization of these organisms.

EXAMPLE 2

Identification and characterization of the *tcf* operon.

The present inventors found that *Salmonella enterica* subspecies I serovar Typhi contains DNA encoding an additional fimbrial operon. the *tcf* operon, in the *sinR-pagN* intergenic region. Southern blot analysis revealed a markedly different restriction pattern in *S. enterica* serovar Typhi than the other subspecies I isolates, suggesting that the *saf-sin* region in serovar Typhi might carry additional DNA relative to serovar Typhimurium strains. A PCR reaction (using a kit from Roche) was therefore performed using a *sinR* (5'-GTA AAT CGC TTA GTC GCC-3') specific forward primer and a *pagN* (5'-TCA ACT CAA CCT TCA GCC-3') specific reverse primer.

This primer pair produced, as expected, a product of 2 kb in serovar Typhimurium from the SARC collection, while from serovar Typhi the product was 10 kb. Thus, the neighboring *sinR* and *pagN* genes in serovar Typhimurium strains are separated by approximately 8 kb in serovar Typhi.

The Typhi specific PCR product was purified, digested partially with *EcoRI* and sub-cloned into pUC18 forming a set of overlapping clones. Sequencing of the clones revealed a putative fimbrial operon designated *tcf* for Typhi Colonizing Factor. Four ORFs, *tcfA,B,C,D*, have been identified with putative proteins having significant homology to CooB (38% identical over 192 aa), CooA (37% identical over 170 aa), CooC (34% identical over 872 aa) and CooD (31% identical over 272 aa), respectively. The Coo proteins are involved in the biosynthesis of the CS1 colonizing factor antigens of enterotoxigenic *E.coli* (Fig. 4) (Froehlich et al., 1994). The peptide of the *tcfB* ORF is also homologous to the CblA major fimbrial subunit protein (45% identical over 154 aa) of the cable

type II pili of the cystic fibrosis-associated *Burkholderia cepacia* (Sajjan et al., 1995). Down-stream of the *tcf*-operon two ORFs were identified with the same transcriptional orientation as the *tcf* genes. The first was designated *tinR* for Typhi insert regulator because it is homologous (33% identical over 144 aa) to AzlB of *Bacillus subtilis*, a member of the Lrp/AsnC family of transcriptional regulators (Belitsky et al., 1997). *tinR* is followed by an ORF (*tioA* for Typhi insert orf) encoding a putative protein of 205 amino acids with no significant homologies to anything in the DDBJ/EMBL/GenBank databases. The above sequence from *Salmonella enterica* serovar Typhi strain RKS 3333 and the *tcf* region of the incomplete genome sequence from serovar Typhi strain CT18 ([http:// www.sanger.ac.uk](http://www.sanger.ac.uk)) are 99% identical over the total length of the investigated region in concordance with the clonal nature of the serovar .

A 2 kb large internal *EcoR* I fragment was used as a probe in a Southern blot of the SARC collection. This blot shows that *Salmonella enterica* subspecies I serovar Typhi (SARC2) is the only strain in the collection possessing DNA hybridizing to this fragment (Fig. 4).

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Claims:

1. Peptide encoded by a nucleotide sequence selected from Sequence Listing No. 1 (SEQ ID NO 1) and Sequence Listing No. 2 (SEQ ID NO 2) for use in medicine.
2. Antibodies directed against a peptide encoded by a nucleotide sequence selected from Sequence Listing No. 1 (SEQ ID NO 1) and Sequence Listing No. 2 (SEQ ID NO 2) for use in medicine.
3. Nucleotide sequence selected from Sequence Listing No. 1 (SEQ ID NO 1) and Sequence Listing No. 2 (SEQ ID NO 2) for use in medicine.
4. A vaccine for the protection against diseases caused by *Salmonella enterica* subspecies I, comprising a peptide encoded by a nucleotide sequence selected from Sequence Listing No. 1 (SEQ ID NO 1), or antibodies directed against said peptide and, optionally, a pharmaceutically acceptable carrier.
5. A vaccine for the protection against diseases caused by *Salmonella enterica* subspecies I serovar Typhi, comprising a peptide encoded by a nucleotide sequence selected from Sequence Listing No. 2 (SEQ ID NO 2), or antibodies directed against said peptide and, optionally, a pharmaceutically acceptable carrier.
6. A nucleic acid vaccine for the protection against diseases caused by *Salmonella enterica* subspecies I, comprising a nucleic acid sequence selected from Sequence Listing No. 1 (SEQ ID NO 1) and, optionally, a pharmaceutically acceptable carrier.
7. A nucleic acid vaccine for the protection against diseases caused by *Salmonella enterica* subspecies I serovar Typhi, comprising a nucleic acid sequence selected from Sequence Listing No. 2 (SEQ ID NO 2) and, optionally, a pharmaceutically acceptable carrier.
8. A vector vaccine for the protection against diseases caused by *Salmonella enterica* subspecies I, comprising a host in which a recombinant vector comprising a nucleic acid sequence selected from Sequence Listing No. 1 (SEQ ID NO 1), has been inserted and, optionally, a pharmaceutically acceptable carrier.

9. A vector vaccine for the protection against diseases caused by *Salmonella enterica* subspecies I serovar Typhi, comprising a host in which a recombinant vector comprising a nucleic acid sequence selected from Sequence Listing No. 2 (SEQ ID NO 2), has been inserted and, optionally, a pharmaceutically acceptable carrier.

5

10. A method for protection against diseases caused by *Salmonella enterica* subspecies I, comprising administering a vaccine according to any of claims 4, 6, and 8.

10 11. A method for protection against diseases caused by *Salmonella enterica* subspecies I serovar Typhi, comprising administering a vaccine according to any of claims 5, 7, and 9.

15 12. Antibodies directed against a peptide encoded by a nucleotide sequence selected from Sequence Listing No. 1 (SEQ ID NO 1) and Sequence Listing No. 2 (SEQ ID NO 2), for use in a diagnostic method.

20 13. Peptide encoded by a nucleotide sequence selected from Sequence Listing No. 1 (SEQ ID NO 1) and Sequence Listing No. 2 (SEQ ID NO 2), for use in a diagnostic method.

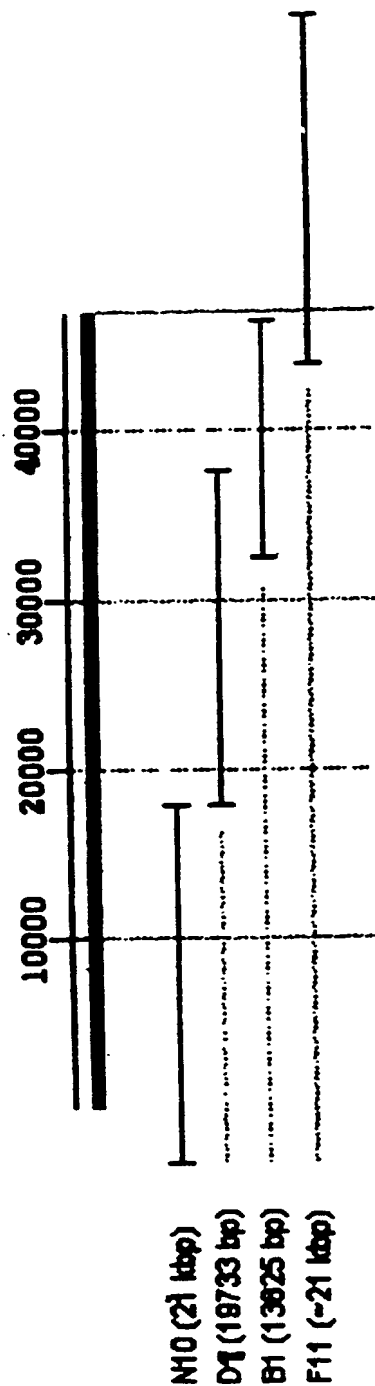
25 14. Primers for, or probes that hybridize with, a nucleotide sequence selected from Sequence Listing No. 1 (SEQ ID NO 1) and Sequence Listing No. 2 (SEQ ID NO 2), for use in a diagnostic method for the purpose of detecting *Salmonella enterica* subspecies I.

ABSTRACT

5 The present invention is based on the finding that two fimbrial structures are specific for *Salmonella enterica* subspecies 1 bacteria. Due to their specificity they can be used to provide vaccines against *Salmonella enterica* subspecies I as well as for detection of *Salmonella enterica* subspecies I.

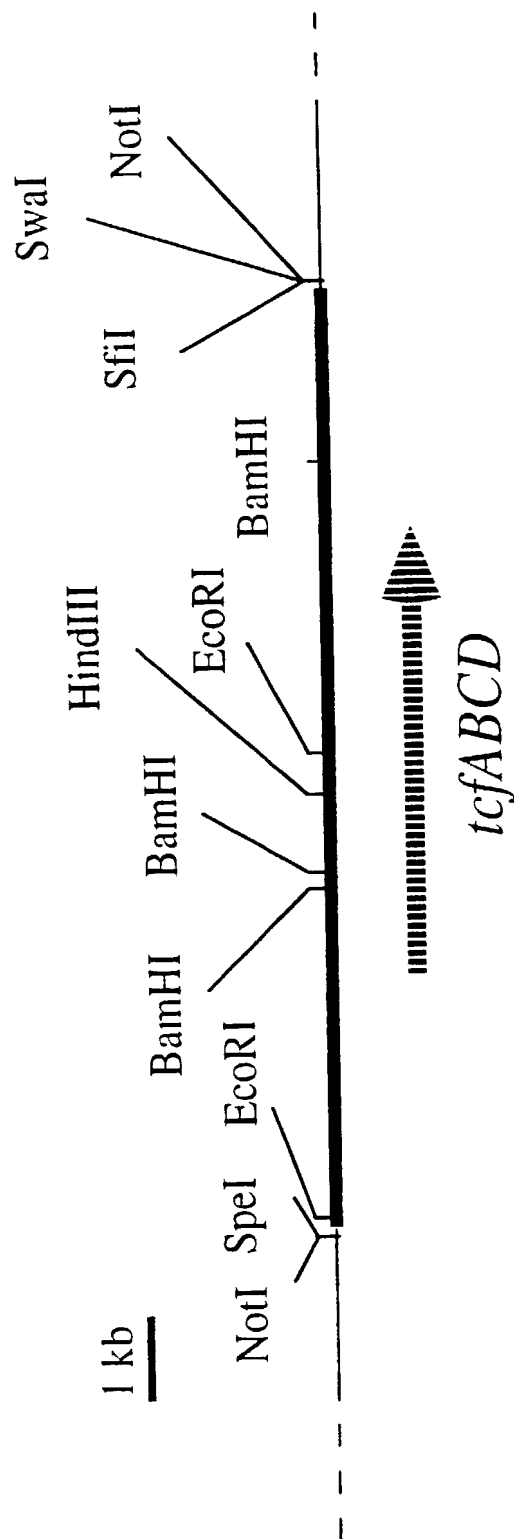
1/4

Fig. 1



2/4

Fig. 2

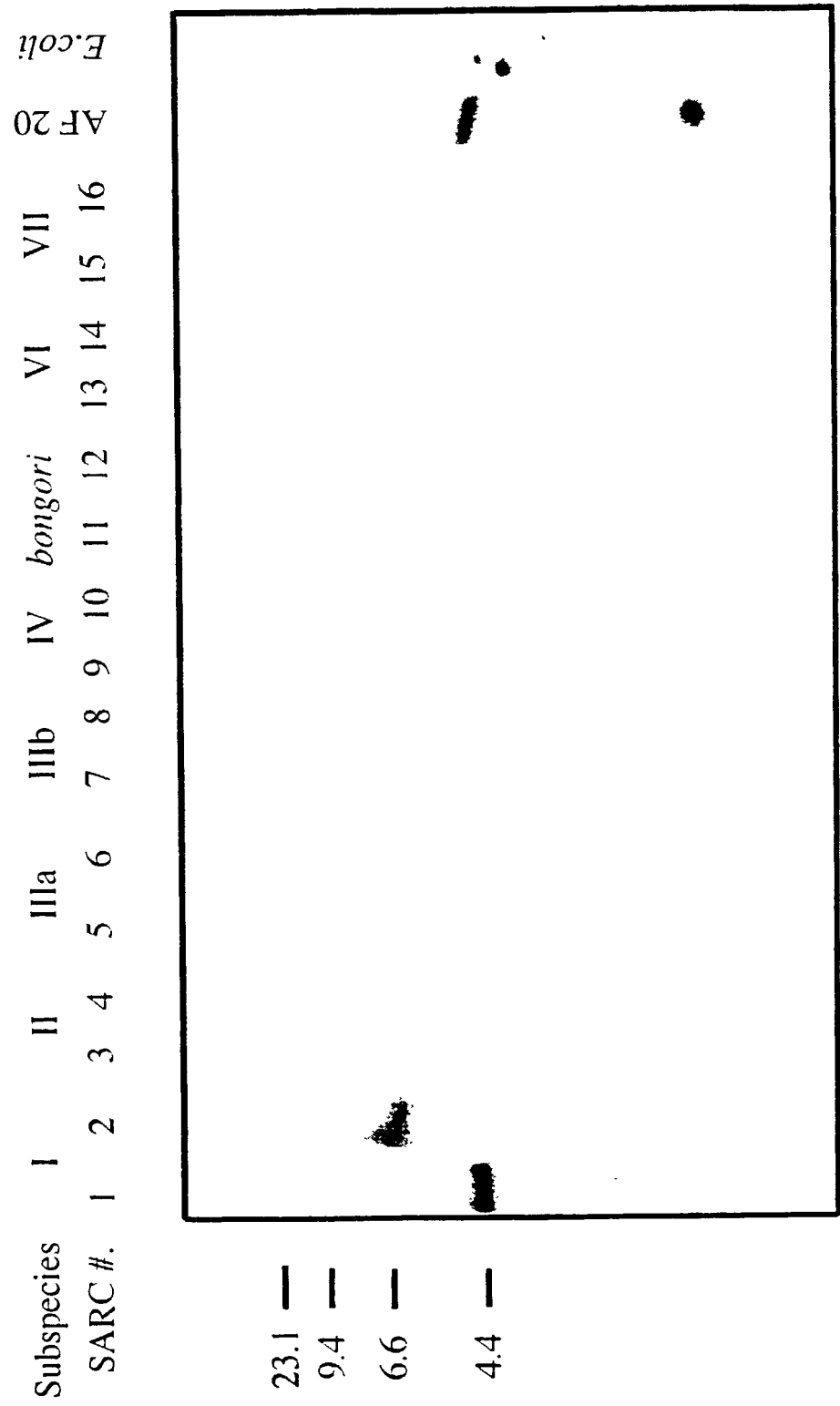


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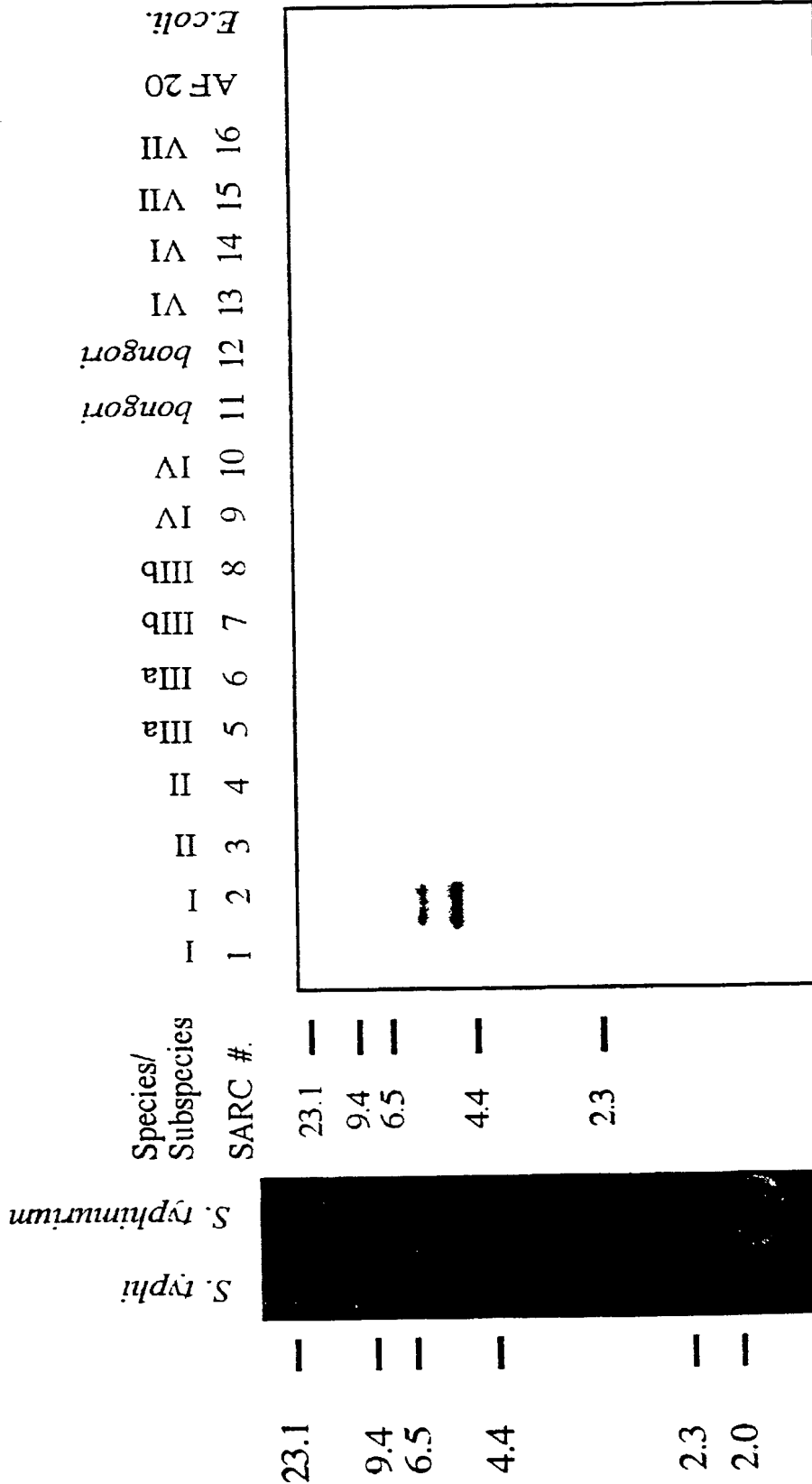
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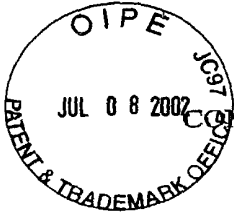
Fig. 3



4/4

Fig. 4





Atty Dkt No. ABR 022 US
PATENT

COMBINED DECLARATION AND POWER OF ATTORNEY
FOR UTILITY PATENT APPLICATION

AS A BELOW-NAMED INVENTOR, I HEREBY DECLARE THAT:

My residence, post office address and citizenship are as stated below next to my name.

I believe I am the original, first and sole inventor (if only one name is listed below) or an original, first and joint inventor (if more than one name is listed below) of the subject matter which is claimed and for which a patent is sought on the invention entitled: FIMBRIAL PROTEINS the specification of which

X was filed on 26 November 2001 and assigned Serial No. 09/926,596; and
X was initially filed in a PCT application designating the US on 26 May 2000, and assigned Serial No. PCT/SE00/01079;
X was amended under PCT Article 19 (35 U.S.C. §371(c)(3)) on 4 September 2001 and on 13 September 2001; and
X has was amended by the Preliminary Amendment dated 22 November 2001.

I HAVE REVIEWED AND UNDERSTAND THE CONTENTS OF THE ABOVE-IDENTIFIED SPECIFICATION, INCLUDING THE CLAIMS, AS AMENDED BY ANY AMENDMENT REFERRED TO ABOVE.

I acknowledge and understand that I am an individual who has a duty to disclose information which is material to the patentability of the claims of this application in accordance with Title 37, Code of Federal Regulations, §§ 1.56(a) and (b) which state:

(a) A patent by its very nature is affected with a public interest. The public interest is best served, and the most effective patent examination occurs when, at the time an application is being examined, the Office is aware of and evaluates the teachings of all information material to patentability. Each individual associated with the filing and prosecution of a patent application has a duty of candor and good faith in dealing with the Office, which includes a duty to disclose to the Office all information known to that individual to be material to patentability as defined in this section. The duty to disclose information exists with respect to each pending claim until the claim is canceled or withdrawn from consideration, or the application becomes abandoned. Information material to the patentability of a claim that is canceled or withdrawn from consideration need not be submitted if the information is not material to the patentability of any claim remaining under consideration in the application. There is no duty to submit information which is not material to the patentability of any existing claim. The duty to disclose all information known to be material to patentability is deemed to be satisfied if all information known to be material to patentability of any claim issued in a patent was cited by the Office or submitted to the Office in the manner prescribed by §§ 1.97(b)-(d) and 1.98. However, no patent will be granted on an application in connection with which

fraud on the Office was practiced or attempted or the duty of disclosure was violated through bad faith or intentional misconduct. The Office encourages applicants to carefully examine:

(1) prior art cited in search reports of a foreign patent office in a counterpart application, and

(2) the closest information over which individuals associated with the filing or prosecution of a patent application believe any pending claim patentably defines, to make sure that any material information contained therein is disclosed to the Office.

(b) Under this section, information is material to patentability when it is not cumulative to information already of record or being made of record in the application, and

(1) It establishes, by itself or in combination with other information, a *prima facie* case of unpatentability of a claim; or

(2) It refutes, or is inconsistent with, a position the applicant takes in:

(i) Opposing an argument of unpatentability relied on by the Office,

or

(ii) Asserting an argument of patentability.

A *prima facie* case of unpatentability is established when the information compels a conclusion that a claim is unpatentable under the preponderance of evidence, burden-of-proof standard, giving each term in the claim its broadest reasonable construction consistent with the specification, and before any consideration is given to evidence which may be submitted in an attempt to establish a contrary conclusion of patentability.

I do not know and do not believe this invention was ever known or used in the United States of America before my or our invention thereof, or patented or described in any printed publication in any country before my or our invention thereof or more than one year prior to said application. This invention was not in public use or on sale in the United States of America more than one year prior to this application. This invention has not been patented or made the subject of an inventor's certificate issued before the date of this application in any country foreign to the United States of America on any application filed by me or my legal representatives or assigns more than six months prior to this application.

I hereby appoint the following attorneys and agents to prosecute that application and to transact all business in the Patent and Trademark Office connected therewith and to file, to prosecute and to transact all business in connection with all patent applications directed to the invention:

3 Thomas P. McCracken, Reg. No. 38,548
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This appointment, including the right to delegate this appointment, shall also apply to the same extent to any proceedings established by the Patent Cooperation Treaty.

I hereby declare that all statements made herein of my own knowledge are true and that all statements made on information and belief are believed to be true; and further that these statements were made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under § 1001 of Title 18 of the United States Code and that such willful false statements may jeopardize the validity of the application or any patent issued thereon.

1-0
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2-0
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3-0
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Gly Gly Lys Pro Ile Thr Gly Leu Glu Tyr Val Pro Pro Phe Ala Asp	
365 370 375	
aaa aca cta aat atg cca ggt agt gcc cat ggt gat atc gag tgg aga	38635
Lys Thr Leu Asn Met Pro Gly Ser Ala His Gly Asp Ile Glu Trp Arg	
380 385 390 395	
gtt att aca gac ttt ggt ggt gaa agt cat ccg ttc cac tac gtt ctt	38683
Val Ile Thr Asp Phe Gly Gly Glu Ser His Pro Phe His Tyr Val Leu	
400 405 410	
aaa taa atccagggggc ttagcggcag aaa atg aag ttc aaa caa cct gcc ttg	38736
Lys Met Lys Phe Lys Gln Pro Ala Leu	
415 420	
cta ctg ttc atc gcg gga gtg gtt cat tgc gca aat gcg cac act tac	38784

Leu	Leu	Phe	Ile	Ala	Gly	Val	Val	His	Cys	Ala	Asn	Ala	His	Thr	Tyr	
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aca	ttc	gat	gca	tca	atg	ttg	ggc	gat	gca	gcg	aaa	ggg	gtt	gat	atg	38832
Thr	Phe	Asp	Ala	Ser	Met	Leu	Gly	Asp	Ala	Ala	Lys	Gly	Val	Asp	Met	
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tcg	ctc	ttt	aac	cag	ggg	tta	caa	cag	cca	ggg	act	tat	cgc	gtg	gac	38880
Ser	Leu	Phe	Asn	Gln	Gly	Leu	Gln	Gln	Pro	Gly	Thr	Tyr	Arg	Val	Asp	
			455					460					465			
gtg	atg	gtg	aat	ggg	aaa	cgt	gtc	gac	acc	cgt	gat	gtg	gtg	ttc	aaa	38928
Val	Met	Val	Asn	Gly	Lys	Arg	Val	Asp	Thr	Arg	Asp	Val	Val	Phe	Lys	
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ttg	gaa	aag	gat	ggg	caa	gga	acg	cct	gtt	ctg	gct	cct	tgt	ttg	acg	38976
Leu	Glu	Lys	Asp	Gly	Gln	Gly	Thr	Pro	Val	Leu	Ala	Pro	Cys	Leu	Thr	
				490					495					500		
gtc	agt	cag	ctt	tca	cgc	tac	ggc	gta	aaa	acg	gaa	gat	tac	cct	cag	39024
Val	Ser	Gln	Leu	Ser	Arg	Tyr	Gly	Val	Lys	Thr	Glu	Asp	Tyr	Pro	Gln	
			505					510					515			
ttg	tgg	aaa	gca	gca	aag	ccc	cca	gat	gag	tgt	gcg	gat	ctg	acc	gcc	39072
Leu	Trp	Lys	Ala	Ala	Lys	Pro	Pro	Asp	Glu	Cys	Ala	Asp	Leu	Thr	Ala	
			520					525					530			
att	cca	cag	gct	aaa	gcg	gta	ctg	gat	atc	aat	aat	cag	caa	ctg	caa	39120
Ile	Pro	Gln	Ala	Lys	Ala	Val	Leu	Asp	Ile	Asn	Asn	Gln	Gln	Leu	Gln	
			535					540				545				
ctg	agt	att	ccg	cag	ttg	gcg	ttg	cgt	ccg	gaa	ttt	aag	ggg	atc	gct	39168
Leu	Ser	Ile	Pro	Gln	Leu	Ala	Leu	Arg	Pro	Glu	Phe	Lys	Gly	Ile	Ala	
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cca	gaa	gat	ctt	tgg	gat	gat	ggg	att	ccg	gcg	ttt	ctg	atg	aac	tac	39216
Pro	Glu	Asp	Leu	Trp	Asp	Asp	Gly	Ile	Pro	Ala	Phe	Leu	Met	Asn	Tyr	
				570					575					580		
agt	gcg	agg	aca	acg	cag	acg	gat	tac	aaa	atg	gat	atg	gtg	ggg	cgt	39264
Ser	Ala	Arg	Thr	Thr	Gln	Thr	Asp	Tyr	Lys	Met	Asp	Met	Val	Gly	Arg	
			585					590					595			
gac	aac	tct	tcc	tgg	gta	caa	ctg	caa	ccg	gga	atc	aat	ata	ggg	gcg	39312
Asp	Asn	Ser	Ser	Trp	Val	Gln	Leu	Gln	Pro	Gly	Ile	Asn	Ile	Gly	Ala	
			600					605				610				
tgg	cgt	gtc	cgc	aat	gcg	acc	agc	tgg	cag	cgg	agt	agt	caa	ctg	tcg	39360
Trp	Arg	Val	Arg	Asn	Ala	Thr	Ser	Trp	Gln	Arg	Ser	Ser	Gln	Leu	Ser	
			615					620				625				
ggg	aag	tgg	cag	gca	gca	tat	acc	tat	gct	gag	cgt	gga	ctg	tac	tca	39408
Gly	Lys	Trp	Gln	Ala	Ala	Tyr										

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ggc aat ctg agt tta acc ggt tcc cgt acc gac tgg cgt aat cgc ccc Gly Asn Leu Ser Leu Thr Gly Ser Arg Thr Asp Trp Arg Asn Arg Pro 920 925 930	40272
ggg cat gat gac agc tac gga ctg agt tgg gga acc tct atc gga ggg Gly His Asp Asp Ser Tyr Gly Leu Ser Trp Gly Thr Ser Ile Gly Gly 935 940 945	40320
ggc tcg ctg tca ttg aac tgg aat caa aac aga acg ctg tgg cgc aat Gly Ser Leu Ser Leu Asn Trp Asn Gln Asn Arg Thr Leu Trp Arg Asn 950 955 960 965	40368
ggc gcg cac cgt aaa gag aac ata acc agc ctg tgg ttc agt atg cca Gly Ala His Arg Lys Glu Asn Ile Thr Ser Leu Trp Phe Ser Met Pro 970 975 980	40416
tta agc cgc tgg acg ggg aat aat gta agt gct agt tgg cag atg act Leu Ser Arg Trp Thr Gly Asn Asn Val Ser Ala Ser Trp Gln Met Thr 985 990 995	40464
tca cca tca cac ggt ggt cag acg caa caa gtg ggg gtc aac gga gag Ser Pro Ser His Gly Gly Gln Thr Gln Gln Val Gly Val Asn Gly Glu 1000 1005 1010	40512
gca ttc agt cag caa ctg gat tgg gag gtg cgt cag agt tac cgt gcc Ala Phe Ser Gln Gln Leu Asp Trp Glu Val Arg Gln Ser Tyr Arg Ala 1015 1020 1025	40560
gat gcc ccg cca ggt ggt ggt aat aac agc gca ttg cac ttg gca tgg Asp Ala Pro Pro Gly Gly Gly Asn Asn Ser Ala Leu His Leu Ala Trp 1030 1035 1040 1045	40608
aat ggg gat tac ggc ctg tta ggt ggt gac tat agc tac agc cgg gcg Asn Gly Asp Tyr Gly Leu Leu Gly Gly Asp Tyr Ser Tyr Ser Arg Ala 1050 1055 1060	40656
atg cgc cag atg gga gtc aat atc gcg gga ggt ata gtt atc cac cat Met Arg Gln Met Gly Val Asn Ile Ala Gly Gly Ile Val Ile His His 1065 1070 1075	40704
cat ggt gtg acg ctg ggg caa cct ttg caa ggc tca gtg gcg ctg gtt His Gly Val Thr Leu Gly Gln Pro Leu Gln Gly Ser Val Ala Leu Val 1080 1085 1090	40752
gaa gcg cca ggg gcc tcg ggg gtg cca gtt ggc ggc tgg cct ggc gtt Glu Ala Pro Gly Ala Ser Gly Val Pro Val Gly Gly Trp Pro Gly Val 1095 1100 1105	40800
aag acg gat ttt cgt ggc gac acc aca gtg ggc aac ctg aac gtc tat Lys Thr Asp Phe Arg Gly Asp Thr Thr Val Gly Asn Leu Asn Val Tyr 1110 1115 1120 1125	40848
cag gag aat aca gtc agc ctc gat ccg tcg cga cta ccg gat gac gca Gln Glu Asn Thr Val Ser Leu Asp Pro Ser Arg Leu Pro Asp Asp Ala 1130 1135 1140	40896
gag gtc aca caa acc gat gtg cgc gtg gtg cca acc gaa ggg gcg gtg Glu Val Thr Gln Thr Asp Val Arg Val Val Pro Thr Glu Gly Ala Val 1145 1150 1155	40944

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gtg gaa gcg aag ttt cac act cgc atc ggg gcc agg gca ctg atg acg Val Glu Ala Lys Phe His Thr Arg Ile Gly Ala Arg Ala Leu Met Thr 1160 1165 1170	40992
ctg aaa cgg gaa gat ggt agc gcc att cct ttc ggg gcg cag gtt aca Leu Lys Arg Glu Asp Gly Ser Ala Ile Pro Phe Gly Ala Gln Val Thr 1175 1180 1185	41040
gtc aat ggg cag gat ggc agt gct gct ctg gtg gat act gat agc cag Val Asn Gly Gln Asp Gly Ser Ala Ala Leu Val Asp Thr Asp Ser Gln 1190 1195 1200 1205	41088
gtt tat ctc act ggt ttg gcg gat aag ggc gaa ctg acg gtg aaa tgg Val Tyr Leu Thr Gly Leu Ala Asp Lys Gly Glu Leu Thr Val Lys Trp 1210 1215 1220	41136
gga gca cag caa tgt cgg gtt aac tac cgc cta cct gcc cac aag gga Gly Ala Gln Gln Cys Arg Val Asn Tyr Arg Leu Pro Ala His Lys Gly 1225 1230 1235	41184
atc gcg ggc ttg tat caa atg agc ggt ctc tgc aga tag ccgattctga Ile Ala Gly Leu Tyr Gln Met Ser Gly Leu Cys Arg 1240 1245 1250	41233
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agc gta agc tta ctg gtc gct gcc agt agc ttg gtg ccg ata gcg aac Ser Val Ser Leu Leu Val Ala Ala Ser Ser Leu Val Pro Ile Ala Asn 1265 1270 1275	41331
gcc gca gaa aaa ctt cag aca acg cta cgt gta ggt act tac ttt cgg Ala Ala Glu Lys Leu Gln Thr Thr Leu Arg Val Gly Thr Tyr Phe Arg 1280 1285 1290 1295	41379
gct ggg cac gtg cca gat ggg atg gtg ctt gcg caa ggc tgg gtg act Ala Gly His Val Pro Asp Gly Met Val Leu Ala Gln Gly Trp Val Thr 1300 1305 1310	41427
tat cac ggc agt cac agc ggg ttt cgg gta tgg agc gat gag caa aag Tyr His Gly Ser His Ser Gly Phe Arg Val Trp Ser Asp Glu Gln Lys 1315 1320 1325	41475
gcg ggt aac acg cct acc gta ttg ctg ctg agc ggg caa cag gat cct Ala Gly Asn Thr Pro Thr Val Leu Leu Leu Ser Gly Gln Gln Asp Pro 1330 1335 1340	41523
cgc cat cac att cag gtt cgc ctg gag ggc gag ggg tgg caa cca gat Arg His His Ile Gln Val Arg Leu Glu Gly Glu Gly Trp Gln Pro Asp 1345 1350 1355	41571
acg gtg agt ggt cgt ggc gcc att tta aga acc gct gca gat aac gcc Thr Val Ser Gly Arg Gly Ala Ile Leu Arg Thr Ala Ala Asp Asn Ala 1360 1365 1370 1375	41619
agt ttc agt gtg gtc gtt gat ggc aat cag gaa gtg cct gcg gac acc Ser Phe Ser Val Val Val Asp Gly Asn Gln Glu Val Pro Ala Asp Thr 1380 1385 1390	41667
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Substitute sheet (Rule 26)

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<210> 2

<211> 166

<212> PRT

<213> Salmonella typhimurium

<400> 2

Met Lys Ser Ile Lys Lys Leu Ile Ile Ala Ser Ala Leu Ser Met Met

31

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Ala Ala Ser Cys Tyr Ala Gly Ser Phe Leu Pro Asn Ser Glu Gln Gln	20	25	30
Lys Ser Val Asp Ile Val Phe Ser Ser Pro Gln Asp Leu Thr Val Ser	35	40	45
Leu Ile Pro Val Ser Gly Leu Lys Ala Gly Lys Asn Ala Pro Ser Ala	50	55	60
Lys Ile Ala Lys Leu Val Val Asn Ser Thr Thr Leu Lys Glu Phe Gly	65	70	75
Val Arg Gly Ile Ser Asn Asn Val Val Asp Ser Thr Gly Thr Ala Trp	85	90	95
Arg Val Ala Gly Lys Asn Thr Gly Lys Glu Ile Gly Val Gly Leu Ser	100	105	110
Ser Asp Ser Leu Arg Arg Ser Asp Ser Thr Glu Lys Trp Asn Gly Val	115	120	125
Asn Trp Met Thr Phe Asn Ser Asn Asp Thr Leu Asp Ile Val Leu Thr	130	135	140
Gly Pro Ala Gln Asn Val Thr Ala Asp Thr Tyr Pro Ile Thr Leu Asp	145	150	155
Val Val Gly Tyr Gln Pro	165		

<210> 3
 <211> 245
 <212> PRT
 <213> Salmonella typhimurium

<400> 3
 Met Lys Ile Val Asn Phe Ala Val Met Ala Val Ala Leu Phe Ala Thr
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 Asn Ser Met Val Ser Val Tyr Ala Val Asn Gln Gln Leu Asn Ser Ala
 20 25 30
 Thr Lys Leu Phe Ser Val Lys Leu Gly Ala Thr Arg Val Ile Tyr His
 35 40 45
 Ala Gly Thr Ala Gly Ala Thr Leu Ser Val Ser Asn Pro Gln Asn Tyr
 50 55 60
 Pro Ile Leu Val Gln Ser Ser Val Lys Ala Ala Asp Lys Ser Ser Pro
 65 70 75 80
 Ala Pro Phe Leu Val Met Pro Pro Leu Phe Arg Leu Glu Ala Asn Gln
 85 90 95
 Gln Ser Gln Leu Arg Ile Val Arg Thr Gly Gly Asp Met Pro Thr Asp
 100 105 110
 Arg Glu Thr Leu Gln Trp Val Cys Ile Lys Ala Val Pro Pro Glu Asn

145	150										155					160		
Ile	Pro	Ala	Phe	Leu	Met	Asn	Tyr	Ser	Ala	Arg	Thr	Thr	Gln	Thr	Asp			
				165					170					175				
Tyr	Lys	Met	Asp	Met	Val	Gly	Arg	Asp	Asn	Ser	Ser	Trp	Val	Gln	Leu			
			180					185					190					
Gln	Pro	Gly	Ile	Asn	Ile	Gly	Ala	Trp	Arg	Val	Arg	Asn	Ala	Thr	Ser			
		195					200					205						
Trp	Gln	Arg	Ser	Ser	Gln	Leu	Ser	Gly	Lys	Trp	Gln	Ala	Ala	Tyr	Thr			
	210					215					220							
Tyr	Ala	Glu	Arg	Gly	Leu	Tyr	Ser	Leu	Lys	Ser	Arg	Leu	Thr	Leu	Gly			
225					230					235					240			
Gln	Lys	Thr	Ser	Gln	Gly	Glu	Ile	Phe	Asp	Ser	Val	Pro	Phe	Thr	Gly			
				245					250					255				
Val	Met	Leu	Ala	Ser	Asp	Asp	Asn	Met	Val	Pro	Tyr	Ser	Glu	Arg	Gln			
			260					265					270					
Phe	Ala	Pro	Val	Val	Arg	Gly	Ile	Ala	Arg	Thr	Gln	Ala	Arg	Val	Glu			
		275					280					285						
Val	Lys	Gln	Asn	Gly	Tyr	Thr	Ile	Tyr	Asn	Thr	Thr	Val	Ala	Pro	Gly			
	290					295					300							
Pro	Phe	Ala	Leu	Arg	Asp	Leu	Ser	Val	Thr	Asp	Ser	Ser	Gly	Asp	Leu			
305					310					315					320			
His	Val	Thr	Val	Trp	Glu	Ala	Asp	Gly	Ser	Thr	Gln	Met	Phe	Val	Val			
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Pro	Tyr	Gln	Thr	Pro	Ala	Ile	Ala	Leu	His	Gln	Gly	Tyr	Leu	Lys	Tyr			
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Ser	Leu	Leu	Ala	Gly	Arg	Tyr	Arg	Ser	Ser	Asp	Ser	Ala	Thr	Asp	Lys			
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Arg	Gln	Ile	Ala	Gln	Ala	Thr	Leu	Met	Tyr	Gly	Leu	Pro	Trp	Asn	Leu			
	370					375					380							
Thr	Ala	Tyr	Gly	Gly	Ile	Gln	Ser	Ala	Thr	His	Asn	Gln	Ala	Ala	Leu			
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Leu	Gly	Leu	Gly	Gly	Ser	Leu	Gly	Arg	Trp	Gly	Ser	Leu	Ser	Val	Asp			
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Gly	Ser	Asp	Thr	His	Ser	Gln	Arg	Gln	Gly	Glu	Ala	Val	Gln	Gln	Gly			
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Ala	Ser	Trp	Arg	Leu	Arg	Tyr	Ser	Asn	Gln	Leu	Thr	Ala	Thr	Gly	Thr			
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Asn	Phe	Phe	Leu	Thr	Arg	Trp	Gln	Tyr	Ala	Ser	Gln	Gly	Tyr	Asn	Thr			
	450					455					460							
Leu	Ser	Asp	Val	Leu	Asp	Ser	Tyr	Arg	His	Asn	Gly	Asn	Arg	Leu	Trp			
465					470					475					480			

Ser	Trp	Arg	Glu	Asn 485	Leu	Gln	Pro	Ser	Ser 490	Arg	Thr	Thr		Leu	Met	Leu 495
Ser	Gln	Ser	Trp 500	Gly	Arg	His	Leu	Gly 505	Asn	Leu	Ser	Leu	Thr 510	Gly	Ser	
Arg	Thr	Asp 515	Trp	Arg	Asn	Arg	Pro 520	Gly	His	Asp	Asp	Ser 525	Tyr	Gly	Leu	
Ser	Trp 530	Gly	Thr	Ser	Ile	Gly 535	Gly	Gly	Ser	Leu	Ser 540	Leu	Asn	Trp	Asn	
Gln 545	Asn	Arg	Thr	Leu	Trp 550	Arg	Asn	Gly	Ala	His 555	Arg	Lys	Glu	Asn	Ile 560	
Thr	Ser	Leu	Trp	Phe 565	Ser	Met	Pro	Leu	Ser 570	Arg	Trp	Thr	Gly	Asn 575	Asn	
Val	Ser	Ala 580	Ser	Trp	Gln	Met	Thr	Ser 585	Pro	Ser	His	Gly	Gly 590	Gln	Thr	
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Glu 610	Val	Arg	Gln	Ser	Tyr	Arg 615	Ala	Asp	Ala	Pro 620	Pro	Gly	Gly	Gly	Asn	
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Ala	Gly	Gly	Ile 660	Val	Ile	His	His	His 665	Gly	Val	Thr	Leu	Gly 670	Gln	Pro	
Leu	Gln	Gly 675	Ser	Val	Ala	Leu	Val 680	Glu	Ala	Pro	Gly	Ala 685	Ser	Gly	Val	
Pro 690	Val	Gly	Gly	Trp	Pro	Gly 695	Val	Lys	Thr	Asp	Phe 700	Arg	Gly	Asp	Thr	
Thr 705	Val	Gly	Asn	Leu	Asn 710	Val	Tyr	Gln	Glu	Asn 715	Thr	Val	Ser	Leu	Asp 720	
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Val	Val	Pro 740	Thr	Glu	Gly	Ala	Val 745	Val	Glu	Ala	Lys	Phe 750	His	Thr	Arg	
Ile	Gly	Ala 755	Arg	Ala	Leu	Met	Thr 760	Leu	Lys	Arg	Glu	Asp 765	Gly	Ser	Ala	
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Lys Gly Glu Leu Thr Val Lys Trp Gly Ala Gln Gln Cys Arg Val Asn
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Val Pro Asp Gly Met Val Leu Ala Gln Gly Trp Val Thr Tyr His Gly
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Ser His Ser Gly Phe Arg Val Trp Ser Asp Glu Gln Lys Ala Gly Asn
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Ile Gln Val Arg Leu Glu Gly Glu Gly Trp Gln Pro Asp Thr Val Ser
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Gly Arg Gly Ala Ile Leu Arg Thr Ala Ala Asp Asn Ala Ser Phe Ser
115 120 125

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SEQUENCE LISTING NO. 2

<110> Folkesson, Anders

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<130> The tcf insert in Salmonella typhi

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Val Gln Lys Asp Ile Thr Val Thr Ala Asn Ile Asp Ser Thr Leu Glu			
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Leu Leu Gln Ala Asp Gly Ser Ser Leu Pro Ser Thr Met Lys Leu Asp			
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Val Thr Leu Gly Gly Arg Ser Leu Thr Thr Thr Asn Ser Val Leu Glu			
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Ala Lys Thr Leu Phe Pro Asp Gly Lys Thr Gly Asp Ala Ser Ala Leu			
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Val Thr Ala Gly			
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Phe Thr Ser Gln Ala Thr Leu Ile Pro Pro Pro Gly Phe Glu Ser Leu			
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aac agc aat ctg gtc tgt ggt gtc tca gaa gca aaa gac agc agc gag Asn Ser Asn Leu Val Cys Gly Val Ser Glu Ala Lys Asp Ser Ser Glu 520 525 530	3674
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cac agt cgt aat atg aca ctg aat ggt agc ggt gcc ctg ggg ctt ggt His Ser Arg Asn Met Thr Leu Asn Gly Ser Gly Ala Leu Gly Leu Gly 600 605 610	3914
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gaa cat tac aat aac agt cag gcc tgg ttt gac aat ctg ttt gtc cgt Glu His Tyr Asn Asn Ser Gln Ala Trp Phe Asp Asn Leu Phe Val Arg 635 640 645	4010
cag gat ctc ggc aat cag tat tat ctc cag gct ggt cgg atg gat cag Gln Asp Leu Gly Asn Gln Tyr Tyr Leu Gln Ala Gly Arg Met Asp Gln 650 655 660	4058
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ccc ctg agc cgg ttt gat gga tta cga acc ggg acc acc caa gct tat Pro Leu Ser Arg Phe Asp Gly Leu Arg Thr Gly Thr Thr Gln Ala Tyr 680 685 690	4154
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Ser	Gln	Phe	Leu	Thr	Pro	Gly	Met	His	Thr	Leu	Asp	Thr	His	Ser	Leu	
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cca	ccg	gga	agc	tat	cct	ctg	gcg	ttg	cgg	gtg	tat	gag	gat	ggg	att	4346
Pro	Pro	Gly	Ser	Tyr	Pro	Leu	Ala	Leu	Arg	Val	Tyr	Glu	Asp	Gly	Ile	
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Leu	Arg	Arg	Thr	Glu	Thr	Gln	Pro	Phe	Ser	Lys	Gly	Gly	Asn	Ser	Phe	
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Ser	Ala	Gln	Thr	Gln	Trp	Phe	Ile	Gln	Gly	Gly	Leu	Glu	Asp	Thr	Gly	
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Asp	Lys	Ala	Ser	His	Tyr	Asp	Gly	Glu	Thr	Val	Met	Ala	Ala	Gly	Phe	
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caa	act	ggg	ctg	cgg	aaa	aat	atc	agt	ctg	acc	gaa	ggc	atc	tct	ctg	4538
Gln	Thr	Gly	Leu	Arg	Lys	Asn	Ile	Ser	Leu	Thr	Glu	Gly	Ile	Ser	Leu	
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Ala	His	Glu	Ala	Trp	Tyr	Ser	Glu	Thr	Arg	Leu	Asn	Ser	Gln	His	Ala	
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Val	Leu	Asp	Gly	Thr	Leu	Asp	Leu	Ser	Ala	Gly	Ile	Leu	His	Gly	Thr	
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Asp	Ser	Thr	Ser	Gly	Asn	Thr	Glu	Gln	Val	Thr	Tyr	Asn	Asp	Gly	Phe	
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Ser	Ala	Ser	Leu	Trp	Arg	Asn	His	Thr	Glu	Ser	Asp	Ala	Cys	Ser	Gly	
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Arg	His	Pro	Gln	Ser	Val	His	Ala	Ser	Met	Thr	Cys	Gln	Thr	Ser	Met	
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Asn	Ala	Ser	Leu	Ser	Val	Ser	Val	Gly	Asn	Trp	Tyr	Ala	Leu	Leu	Gly	
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Tyr Phe Leu Thr Pro Gly His Leu Leu Val His Asn Ile Ser Ala Ser				
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Met Ser Arg Leu Tyr Val Gly Arg Val Leu Asp Lys Asp Gly Arg Pro				
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ctg ctg gac gca cag cca ctg aac tat cca ttt ttg tcg ttg gga cct				5786
Leu Leu Asp Ala Gln Pro Leu Asn Tyr Pro Phe Leu Ser Leu Gly Pro				
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Ser Gly Arg Phe Ser Leu Gln Ser Glu His Lys Glu Ser Ser Leu Trp				
1240	1245	1250		
ctg ctg tct aaa aac agg atc ctg cgt tgt ccg atg tca gta cat aaa				5882
Leu Leu Ser Lys Asn Arg Ile Leu Arg Cys Pro Met Ser Val His Lys				
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Arg Arg Asp Val Met Gln Val Val Gly Asp Val Arg Cys Glu Leu Ser				
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Asp Val Asp Ala Leu Pro Gln Ala Leu Gln Ile Ser Pro Arg Val Ile				
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Met Ser Asn Lys Met Lys Trp Thr Ser				
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atg aca gcc cat tgg tca gca att att aat ttc atc cga aaa tat gtt				6126
Met Thr Ala His Trp Ser Ala Ile Ile Asn Phe Ile Arg Lys Tyr Val				
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tat cca gca agg ata att gcc atc ctg ctg atg gct ggc gct aca ctg				6174
Tyr Pro Ala Arg Ile Ile Ala Ile Leu Leu Met Ala Gly Ala Thr Leu				
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Pro Gln Val Ala Asp Ala Ile Thr Val Asp Leu Asn Tyr Asp Lys Asn				
1365	1370	1375		
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Asn Val Ala Val Ile Thr Pro Val Trp Ser Gln Glu Trp Ser Val Ala				
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aat gtg ttg ggg gga tgg gta tgt cgt tca aac agg aat gaa aat gag				6318
Asn Val Leu Gly Gly Trp Val Cys Arg Ser Asn Arg Asn Glu Asn Glu				
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Gly Ala Cys Glu Glu Thr His Leu Val Trp Trp Tyr Ala Phe Gly Ala				

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Leu Arg Ile Pro Ser Glu Glu Leu Ala Lys Ile Pro Thr Ser Gly Thr			
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Trp Lys Ala Thr Leu Val Leu Asp Tyr Leu Gln Trp Gly Gly Asp Asp			
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Pro Arg Val Asp Leu Asn Leu His Arg Met Asn Ala Ser Gln Met Ser			
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Gly Arg Ala Asn Leu Asp Met Cys Leu Tyr Asp Gly Gly Val Lys Ala			
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Arg Ser Leu Gln Met Lys Ile Glu Gly Ser Asn Lys Ser Gly Thr Gly			
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Phe Gln Val Ile Lys Ser Asp Ser Ala Asp Thr Ile Asp Tyr Ala Val			
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Ser Met Asn Tyr Gly Gly Arg Ser Ile Pro Val Thr Arg Gly Val Glu			
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<213> Salmonella typhi

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Arg Ser Leu Thr Thr Thr Asn Ser Val Leu Glu Ala Lys Thr Leu Phe
130 135 140
Pro Asp Gly Lys Thr Gly Asp Ala Ser Ala Leu Leu Asn Leu Asp Ile
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Val Lys Pro Asp Thr Val Gln Phe Met Ser Pro Leu Met Val Leu Glu
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Ser Ser Gly Leu Ala Ala Leu Pro Ala Ala Glu Arg Gln Lys Ala Leu
65 70 75 80
Ala Ala Leu Ser Arg Pro Leu Leu Arg Asn Ser Asn Leu Val Cys Gly
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Val Ser Glu Ala Lys Asp Ser Ser Glu Cys Gly Tyr Val Ala Thr Asp
100 105 110
Lys Glu Asp Val Ala Val Ile Phe Asp Glu Asn Asn Ala Gln Leu Ser
115 120 125
Leu Phe Leu Asn Arg Asp Trp Leu Pro Asp Glu Glu Arg Arg Asp Lys
130 135 140
Arg Trp Leu Thr Pro Thr Pro Glu Gly Val Ser Ala Phe Ile His Arg
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Gln Thr Leu Tyr Leu Ser Asp Asp Leu His Ser Arg Asn Met Thr Leu
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Tyr	Leu	Gln	Ala	Gly	Arg	Met	Asp	Gln	Arg	Asn	Leu	Ser	Ser	Ala	Thr
			225		230					235					240
Gly	Gly	Asp	Phe	Gly	Phe	Ser	Leu	Leu	Pro	Leu	Ser	Arg	Phe	Asp	Gly
				245					250					255	
Leu	Arg	Thr	Gly	Thr	Thr	Gln	Ala	Tyr	Val	Asn	His	Glu	Val	Asp	His
			260					265					270		
Asn	Ala	Thr	Pro	Val	Met	Val	Gln	Val	Thr	Arg	Asn	Ala	Arg	Ile	Asp
			275				280					285			
Ile	Tyr	Arg	Gly	Ser	Glu	Leu	Leu	Gly	Ser	Gln	Phe	Leu	Thr	Pro	Gly
			290			295					300				
Met	His	Thr	Leu	Asp	Thr	His	Ser	Leu	Pro	Pro	Gly	Ser	Tyr	Pro	Leu
			305		310					315					320
Ala	Leu	Arg	Val	Tyr	Glu	Asp	Gly	Ile	Leu	Arg	Arg	Thr	Glu	Thr	Gln
				325					330					335	
Pro	Phe	Ser	Lys	Gly	Gly	Asn	Ser	Phe	Ser	Ala	Gln	Thr	Gln	Trp	Phe
			340					345					350		
Ile	Gln	Gly	Gly	Leu	Glu	Asp	Thr	Gly	Asp	Lys	Ala	Ser	His	Tyr	Asp
			355				360					365			
Gly	Glu	Thr	Val	Met	Ala	Ala	Gly	Phe	Gln	Thr	Gly	Leu	Arg	Lys	Asn
			370			375					380				
Ile	Ser	Leu	Thr	Glu	Gly	Ile	Ser	Leu	Ala	His	Glu	Ala	Trp	Tyr	Ser
			385		390					395					400
Glu	Thr	Arg	Leu	Asn	Ser	Gln	His	Ala	Val	Leu	Asp	Gly	Thr	Leu	Asp
				405					410					415	
Leu	Ser	Ala	Gly	Ile	Leu	His	Gly	Thr	Asp	Ser	Thr	Ser	Gly	Asn	Thr
			420				425						430		
Glu	Gln	Val	Thr	Tyr	Asn	Asp	Gly	Phe	Ser	Ala	Ser	Leu	Trp	Arg	Asn
			435			440						445			
His	Thr	Glu	Ser	Asp	Ala	Cys	Ser	Gly	Arg	His	Pro	Gln	Ser	Val	His
			450			455					460				
Ala	Ser	Met	Thr	Cys	Gln	Thr	Ser	Met	Asn	Ala	Ser	Leu	Ser	Val	Ser
			465		470					475					480
Val	Gly	Asn	Trp	Tyr	Ala	Leu	Leu	Gly	Tyr	Ser	Thr	Ser	Arg	Thr	Glu
				485					490					495	

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Ser Glu His Lys Glu Ser Ser Leu Trp Leu Leu Ser Lys Asn Arg Ile
820 825 830
Leu Arg Cys Pro Met Ser Val His Lys Arg Arg Asp Val Met Gln Val
835 840 845
Val Gly Asp Val Arg Cys Glu Leu Ser Asp Val Asp Ala Leu Pro Gln
850 855 860
Ala Leu Gln Ile Ser Pro Arg Val Ile Arg Leu Leu Asn Val Ala Gly
865 870 875 880
Leu Leu Arg His Ser Val Gln Glu Ala
885

<210> 5
<211> 359
<212> PRT
<213> Salmonella typhi

<400> 5
Met Ser Asn Lys Met Lys Trp Thr Ser Met Thr Ala His Trp Ser Ala
1 5 10 15
Ile Ile Asn Phe Ile Arg Lys Tyr Val Tyr Pro Ala Arg Ile Ile Ala
20 25 30
Ile Leu Leu Met Ala Gly Ala Thr Leu Pro Gln Val Ala Asp Ala Ile
35 40 45
Thr Val Asp Leu Asn Tyr Asp Lys Asn Asn Val Ala Val Ile Thr Pro
50 55 60
Val Trp Ser Gln Glu Trp Ser Val Ala Asn Val Leu Gly Gly Trp Val
65 70 75 80
Cys Arg Ser Asn Arg Asn Glu Asn Glu Gly Ala Cys Glu Glu Thr His
85 90 95
Leu Val Trp Trp Tyr Ala Phe Gly Ala Tyr Ser Lys Ile Arg Leu Arg
100 105 110
Phe Arg Glu Gln Ile Ser His Ala Glu Ile Thr Leu Ile Leu Leu Gly
115 120 125
Ser Val Arg Asp Ala Cys Tyr Thr Gly Val Ile Asn Met Asn Ala Ala
130 135 140
Ala Cys Gln Trp Gly Arg Ser Leu Lys Leu Arg Ile Pro Ser Glu Glu
145 150 155 160
Leu Ala Lys Ile Pro Thr Ser Gly Thr Trp Lys Ala Thr Leu Val Leu
165 170 175
Asp Tyr Leu Gln Trp Gly Gly Asp Asp Pro Leu Gly Thr Ser Thr Thr
180 185 190
Asp Ile Thr Leu Asn Val Thr Asp His Phe Ala Glu Asn Ala Ala Ile
195 200 205

Tyr Phe Pro Gln Phe Gly Thr Ala Thr Pro Arg Val Asp Leu Asn Leu
 210 215 220
 His Arg Met Asn Ala Ser Gln Met Ser Gly Arg Ala Asn Leu Asp Met
 225 230 235 240
 Cys Leu Tyr Asp Gly Gly Val Lys Ala Arg Ser Leu Gln Met Lys Ile
 245 250 255
 Glu Gly Ser Asn Lys Ser Gly Thr Gly Phe Gln Val Ile Lys Ser Asp
 260 265 270
 Ser Ala Asp Thr Ile Asp Tyr Ala Val Ser Met Asn Tyr Gly Gly Arg
 275 280 285
 Ser Ile Pro Val Thr Arg Gly Val Glu Phe Ser Leu Asp Asn Val Asp
 290 295 300
 Lys Ala Ala Thr Arg Pro Val Val Leu Pro Gly Gln Arg Gln Ala Val
 305 310 315 320
 Arg Cys Val Pro Val Pro Leu Thr Leu Thr Thr Gln Pro Phe Asn Ile
 325 330 335
 Arg Glu Lys Arg Ser Gly Glu Tyr Gln Gly Thr Leu Thr Val Thr Met
 340 345 350
 Leu Met Gly Thr Gln Thr Pro
 355

<210> 6
 <211> 151
 <212> PRT
 <213> Salmonella typhi

<400> 6
 Met Thr Val Tyr Leu Asp Asp Lys Asp Lys Glu Leu Leu Lys Glu Ile
 1 5 10 15
 Gln Lys Asp Cys Ala Gln Thr Leu Trp Gln Leu Ala Tyr Lys Val Gly
 20 25 30
 Leu Thr Pro Thr Pro Cys Phe Lys Arg Leu Lys Lys Leu Lys Asp Arg
 35 40 45
 Gly Val Ile Ile Gly Gln Phe Ala Leu Leu Asp Lys Glu Lys Leu Gly
 50 55 60
 Leu Ser Leu Asn Val Phe Ile Met Ile Asn Ile Ser Glu Glu Gln Tyr
 65 70 75 80
 Ala Ser Ile Ser Glu Lys Ile Lys Ser Met Pro Glu Val Ile Ala Phe
 85 90 95
 Tyr Arg Ile Ser Gly Ser Phe Asn Tyr Leu Met His Thr Val Phe Thr
 100 105 110

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Asp Met Asn Asp Tyr Tyr Ser Phe Tyr Glu Lys Ile Ile Leu Thr Asn
 115 120 125
 Ser Ser Ile Ser Gly Ser Ala Ser Ser Phe Val Leu Glu Gln Ile Lys
 130 135 140
 Glu Thr Asn Glu Leu Ser Val
 145 150